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(71) Applicant: SEIKO INSTRUMENTS INC.
31-1, Kameido 6-chome
Koto-ku
Tokyo 136(JP)
Applicant: SEIKO ELECTRONIC COMPONENTS
LTD.

3-1, Nishitaga 5-chome,
Taihaku-ku
Sendai-shi Miyagi(JP)

(72) Inventor: Watanabe, Shunji, c/o Seiko
Instruments Inc.
31-1, Kameido 6-chome, Koto-ku
Tokyo(JP)
Inventor: Sakai, Tsugio, c/o Seiko Electronic
Components Ltd
3-1, Nishitaga 5-chome,
Taihaku-ku
Sendai-shi, Miyagi(JP)
Inventor: Sakamoto Hideo c/o Seiko
Electronic Components Ltd
3-1, Nishitaga 5-chome,
Taihaku-ku
Sendai-shi, Miyagi(JP)

(74) Representative: Fleuchaus, Leo, Dipl.-Ing. et al
Melchiorstrasse 42
D-81479 München (DE)

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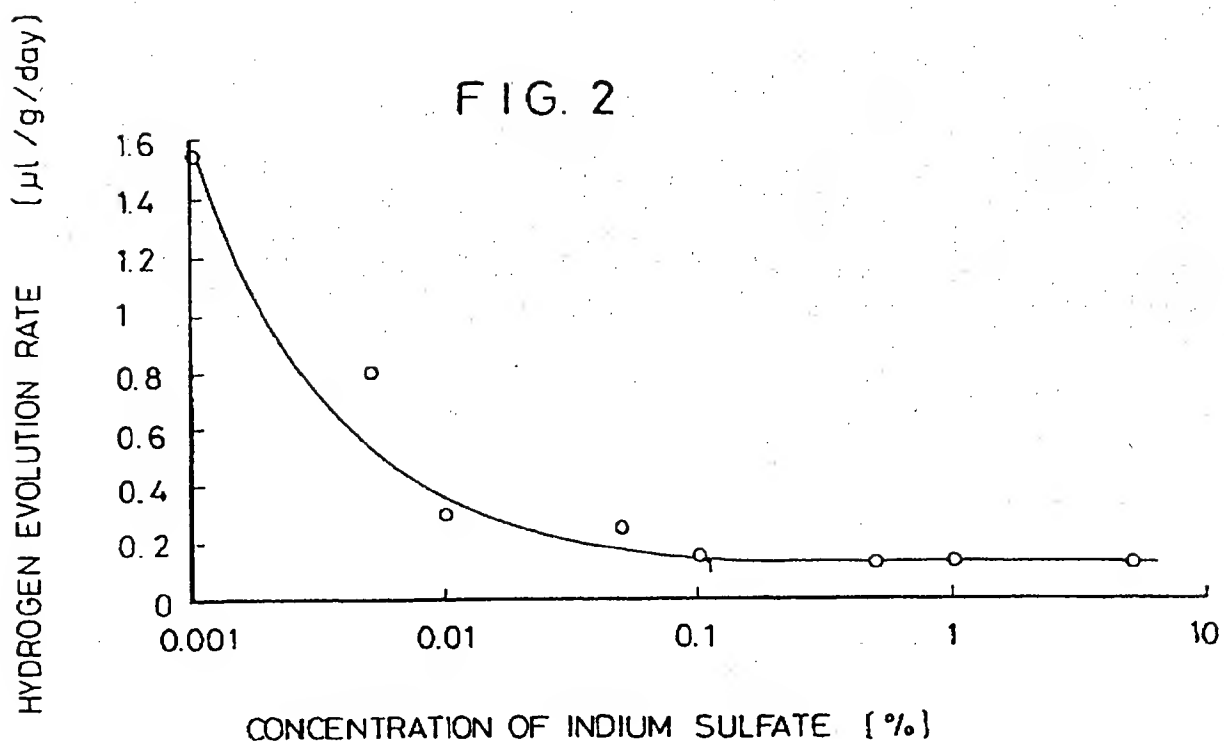
(57) This invention relates to an alkaline battery in which hydrogen gas evolution and deterioration of battery performance can be prevented by using a collector having an outermost layer coated with zinc or metal having a higher hydrogen overvoltage than that of zinc and adding an inhibitor selected from an indium compound, lead oxide, a hydroxide of alkaline earth metal and polyoxyethylen alkylamide to an electrolyte solution or a negative electrode active material.

According to the present invention, it is possible to manufacture an alkaline battery in which evolution of hydrogen can be restrained without involving deterioration of battery performances. Especially, the present invention is effective in a coin type and a button type battery which is sensitive to the evolution of gas and is required to have high performance.

Further, the inventive collector which is plated with zinc alloy more than 3 μ m in thickness and used for low drain such as watches can attain a capacity equivalent to or more than that of a battery using amalgamated zinc. Further, watches loaded with the inventive alkaline batteries which are discarded do not pollute the environment, as they do not include mercury to be a pollution material.

EP 0 582 293 A2

FIG. 2



BACKGROUND OF THE INVENTION

The present invention relates to an alkaline battery which uses mercuryless zinc powder as a negative electrode active material, and uses silver oxide, manganese dioxide, oxygen or the like as a positive electrode active material, and a clock or watch which uses the alkaline battery.

There has been such a task that zinc powder used in the alkaline battery is corroded and dissolved by an electrolyte solution which is an alkali, and the generation of hydrogen gas and the self-discharge in battery performance accompanying therewith are large. In addition, a battery is formed by a collector such as copper or the like which contacts with zinc, and hydrogen gas has been also generated therefrom. In the prior art, as a countermeasure to prevent the above, it has been performed that zinc is mercurated with mercury having a high hydrogen overvoltage, or zinc oxide is added to an electrolyte solution up to approximate saturation.

However, environmental pollution due to mercury from used dry batteries has become worried in recent years, and various studies for realization of reducing mercury or to be mercury legs have been performed. Alloy formation from zinc, plating of a collector, and addition of an organic or inorganic inhibitor to an electrolyte solution fall thereunder.

The alloy formation of zinc has been performed for a fairly long time, and metals such as bismuth, indium, lead and the like have been investigated. Many patent applications have been also filed, and for example, there are Japanese Patent Publication No. 25-27822 (1950), Japanese Patent Publication No. 33-3204 (1958), Japanese Patent Publication No. 63-3942 (1988), Japanese Patent Application Laid-open No. 1-10861 (1989) and the like.

As the inorganic inhibitor, indium oxide and indium hydroxide as the indium compound have been frequently studied, and many patent applications have been also filed.

For example, Japanese Patent Publication No. 51-36450 (1976), Japanese Patent Application Laid-open No. 49-93831 (1974), Japanese Patent Application Laid-open No. 49-112125 (1974), Japanese Patent Application Laid-open No. 59-186255, Japanese Patent Application Laid-open No. 59-186256 (1984), and Japanese Patent Application Laid-open No. 4-26061 (1992) fall thereunder. As those in which a compound of alkaline earth metal is used, there are Japanese Patent Application Laid-open No. 49-8727 (1974), Japanese Patent Application Laid-open No. 49-93831 (1974), Japanese Patent Application Laid-open No. 49-121926 (1974) and the like. As those for organic inhibitors, there are Japanese Patent Application Laid-open No. 2-86064 (1990), Japanese Patent Application Laid-open No. 3-29270 (1991) and the like.

On the other hand, with respect to the collector, it has been performed that the surface is coated with indium or tin having a high hydrogen overvoltage by means of a method of plating or the like, and the formation of a battery due to the contact with zinc is prevented so as to suppress the generation of hydrogen. As patents, there are Japanese Patent Application Laid-open No. 52-74834 (1977), Japanese Patent Laid-open No. 52-98929, Japanese Patent Application Laid-open No. 60-221958 (1985), Japanese Patent Application Publication No. 52-42211 (1977) and the like.

In the prior art, the investigation has been made as individual techniques as described above, however, there has been a strong anticorrosion agent being the mercury, so that optimization by combination of the characteristics of each technique by understanding them has not been frequently performed.

Mercury, which is added in order to prevent corrosion and dissolution of zinc, is not only expensive from a viewpoint of cost, but also includes a large problem of environmental pollution. In addition, the addition of zinc oxide also includes such a task that the viscosity of the electrolyte is raised, and the conductivity is lowered.

Indium oxide and indium hydroxide as the inorganic inhibitor also includes many problems.

Indium oxide is extremely difficult to dissolve in the electrolyte solution which is a caustic alkali, and hydrogen gas is consequently generated due to the contact of indium oxide and with zinc powder or the collector. This is considered to be due to the fact that the solubility of indium oxide is poor, it is impossible to supply indium ion of a degree to sufficiently coat the zinc surface and the collector surface, and indium oxide led to have conductivity due to inevitable impurities during production contacts with zinc and the collector, resulting in formation of a local battery.

It is said that as compared with indium oxide, indium hydroxide to some extent dissolves in an electrolyte solution of caustic alkali, and its solubility relates to the size and crystallinity of particles, however, as compared with indium compounds such as indium sulfate, indium sulfamate, indium chloride and the like, it is extremely difficult to dissolve. Thus, the same task as that of indium oxide occurs. In addition, indium as an amphoteric compound generates polyion together with hydroxide ion (those similar to the description in Inorganic Chemistry Series 7, Coordinate Stereochemistry, written by Yoichi NIIMURA, published by Baihukan Co. Ltd., 65-66), and increases the viscosity of the electrolyte solution, so that it

lowers the conductivity of the electrolyte solution and deteriorates battery performance.

Considerable effects are appreciated in the use of the indium compound easy to dissolve in the electrolyte solution as the inhibitor as compared with the case in which the conventional scarcely soluble inhibitor is used. However, in order to further utilize the characteristic of the indium compound, it is necessary to also solve problems as follows.

The electrode potential of zinc is lower than the deposition potential of indium, so that when indium ion is present in the electrolyte, indium is deposited as metal on zinc and the collector contacting with zinc. However, hydrogen generation is accompanied as a competitive reaction in accordance with the deposition reaction of indium, so that it has been a cause of occurrence of deficiency such as liquid leakage and expansion of the alkaline battery. In addition, there has been such a problem that indium ion not deposited precipitates as hydroxide and decreases the conductivity of the electrolyte solution.

Other than the indium compound, compounds of metals having a relatively high hydrogen overvoltage such as tin and lead are used as the inhibitor, however, there have been problems as follows.

Metal ion, which is supplied from the metal compounds of these metals to the electrolyte solution, is reduced on the surface of zinc and the collector, and deposited as metal. However, when the surface is coated with one species of metal, crystal grains become coarse, it is impossible to homogeneously coat the surface, and the effect is reduced to the half. It is difficult to suppress the hydrogen generation and improve the discharge characteristic by means of single metal. In addition, the compounds of indium and the like are expensive, so that the use by only one species becomes expensive from a viewpoint of cost.

With respect to the corrosion and dissolution of zinc, there are considered a case in which zinc itself is corroded by water and hydroxyl group in the alkaline solution, and a case in which a local battery is formed by the contact with metals such as copper, brass and the like of the collector which are nobler than zinc resulting in dissolution. Thus, such an attempt has been most frequently made that a metal having a high hydrogen overvoltage is added to zinc to form an alloy so as to suppress corrosion and dissolution. It is known that the effect thereof is remarkably expressed especially when indium is added. When zinc is used in which indium is added in a relatively high concentration by, for example, not less than 400 ppm, a part of indium and zinc is once dissolved by the contact with copper and the like of the collector. It is considered that the corrosion and dissolution of zinc are suppressed by a mechanism that the dissolved indium ion is then deposited on the collector, an a film of indium is formed on the collector. However, there has been such a problem that the amount of indium ion to be reduced on the collector is extremely small in the initial state of the contact between zinc and the collector, so that hydrogen is reduced, and the hydrogen gas is generated.

Such an attempt has been most frequently made that the corrosion and dissolution are intended to be suppressed for zinc and the collector using the inhibitor which is the compound of metal nobler than zinc. However, in the case of conventional zinc in which there are much impurities such as iron, it is necessary to use a large amount of the inhibitor, and it has been impossible to suppress the corrosion and dissolution of zinc unless, for example, a large amount of lead monoxide which is a non-pharmaceutical harmful substance and may be an environmental pollution substance is used. Further, there has also been such a problem that when large amounts of lead monoxide and the indium compound are added, needle-like crystals are deposited, which break through a separator to cause a short circuit.

Next, when the task is considered by focusing to a coin or button type silver oxide battery among the alkaline batteries, it is not too much to say that mercurated zinc has been used in the coin or button type silver oxide battery, so that it has been escaped from the task such as the hydrogen gas generation, self-discharge and the like.

In recent years, owing to efforts for improvement in an additive to zinc powder, a separator, a sealing agent and the like, change in a gelling agent and the like, the realization of no mercury has been made for cylindrical alkaline batteries. However, in the case of the coin or button type silver oxide battery in which there is no escape place for hydrogen gas due to the structural reason, there are tasks such as occurrence of expansion and liquid leakage due to gas pressure, self-discharge and the like, and the realization of no mercury has not been achieved.

Until now, there have been mentioned the problems in the conventional inhibitor and the water amount in the battery. Using the improvement in them, the alkaline battery characteristics can be considerably improved. However, it is not achieved to fill a flaw of removal of mercury. Further, tasks to achieve the improvement will be described hereinafter.

When only the conventional inorganic inhibitor is used, there have been such problems that the inhibitor is not homogeneously distributed on the collector because of a fairly small amount of the electrolyte solution in an actual battery, and no metal coating is given, or if there are bubbles between a negative electrode combined agent and the collector, such a place is also generated only in which no metal coating

is made at all.

It is a fairly effective means to manufacture the collector using metal such as indium and tin, or to plate with these metals.

However, when tin is used, it is possible to suppress the hydrogen gas generation as compared with a case in which a collector of copper is used, however, no effect of a degree equivalent to the case of the use of mercury has been obtained.

When indium is used, although the effect is certainly larger than that of tin, there has been such a problem that the raw material is expensive, and the cost increases. In the case of indium plating, there have been such problems that application is poor, no homogeneous film is provided, impurities remain on the surface, and the effect is weakened.

In addition, when no inhibitor is used only with a collector coated with metal, there has been such a problem that a countermeasure is insufficient for the hydrogen gas formation in the case of storage on the way of discharge.

Further, it has been impossible to ensure sufficient prevention from decrease in self-discharge even when fluorocarbon/polyoxyethylene series, polyoxyethylene alkylamide and the like are used which are organic inhibitors considered as effective in the prior art.

Namely, it has been found that there is such a problem that no sufficient effect is obtained using each of the corrosion preventing methods in the prior art. As a result of reconsideration of the role of each inhibitor, it has been found out that better effects are expressed by using a collector in which zinc or metal having a hydrogen overvoltage higher than that of zinc is coated, together with various inhibitors.

When a higher capacity is aimed at, there is such a task that the decrease in the battery capacity due to the absence of mercury must be compensated. In the case of the cylindrical alkaline dry battery, it is accomplished to increase zinc powder as the active material, however, there has been such a problem that in the case of the button type or coin type alkaline battery in which there is no room from a viewpoint of space, such accomplishment is almost impossible.

On the other hand, as viewed from a viewpoint of a clock or watch on which an alkaline battery containing mercury is carried, the battery containing mercury is recovered at retail stores in the case of battery exchange, however, there has been such a problem that when the main body of the clock or watch is discarded, the mercury as an environmental pollution substance is also discarded. Especially, in accordance with the proceeding of low price of the clock or watch, there is such a possibility that the number of clocks or watches to be discarded due to service life expiration of batteries increases.

SUMMARY OF THE INVENTION

An object of the present invention is to provide an alkaline battery which does not contain mercury.

Another object of the present invention is to provide an alkaline battery comprising mercuryless zinc powder as a negative electrode active material.

Still another object of the present invention is to provide an applied product using an alkaline battery which does not contain mercury.

A further object of the present invention is to provide an applied product using an alkaline battery comprising mercuryless zinc powder as a negative electrode active material.

When an indium compound such as indium sulfate, indium sulfamate, indium chloride or the like is added as an inhibitor into an electrolyte solution or a negative electrode active material, so as to allow indium ion to exist in the electrolyte solution in an amount sufficient to coat zinc and the collector, indium can be immediately deposited on zinc and the negative electrode collector. By coating zinc and the collector with indium having a high hydrogen overvoltage, it is possible to prevent corrosion and dissolution of them.

Further, in order to effectively utilize these inhibitors, a complexing agent is added into the electrolyte solution beforehand, and indium ion which is generated by dissolution of the indium compound is subjected to complexing, thereby it is possible to solve the tasks of the hydrogen gas formation during deposition of indium and the decrease in the conductivity of the electrolyte solution due to precipitation of not deposited indium as hydroxide.

The problem caused by the use of the inhibitor of only one species of the compound of metal can be solved by adding two or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin and lead oxide into the electrolyte solution or the negative electrode active material, and depositing metal contained in these compounds as an alloy onto zinc and the negative electrode collector. Characteristics, which cannot be obtained by single metal coating, are obtained by depositing the two or more species of metals.

When zinc in which indium is added in a relatively high concentration is used, it is considered that [the (corrosion and dissolution) of (zinc or the collector)] are suppressed owing to a mechanism that indium ion dissolved from zinc is deposited on the collector, and a film of indium is formed on the collector. However, in the initial stage of the contact between zinc and the collector, the amount of indium ion to be reduced on the collector is extremely small. Thus, when the one or more species of the compounds selected from the indium compound which is nobler than zinc, the tin compound containing tetravalent tin and lead oxide are added as the inhibitor into the electrolyte solution or the negative electrode active material, and the metal ion is allowed to exist in the electrolyte solution in an amount sufficient to coat zinc and the collector, then it is possible to immediately deposit lead onto zinc and the negative electrode collector, and it is possible to suppress the hydrogen generation. In this case, depending on the amount of indium in the zinc powder, it is preferable that the adding amount of the inhibitor is about 10-1000 ppm with respect to the zinc powder. If the adding amount is small, it is impossible to sufficiently coat zinc and the collector, while the adding amount is too large, there is such a bad effect that needle-like crystals penetrate through a separator to cause a short circuit.

When mercuryless zinc powder in which the content of iron is not more than 4 ppm with respect to a weight of zinc is used, one or more species of compounds selected from an indium compound nobler than zinc, a tin compound containing tetravalent tin and lead oxide are added as the inhibitor into the electrolyte solution or the negative electrode active material, and ion is allowed to exist in the electrolyte solution in an amount sufficient to coat zinc and the collector, then it is possible to immediately deposit a coating film of metal which is nobler than zinc on zinc and the negative electrode collector, and it is possible to suppress the corrosion and dissolution of zinc and the hydrogen generation accompanying therewith. In this case, it is preferable that the adding amount of the inhibitor is about 10-1000 ppm with respect to the zinc powder. If the adding amount is small, it is impossible to sufficiently coat zinc and the collector, and there is such a bad effect that needle-like crystals penetrate through a separator to cause a short circuit.

Using mercuryless zinc containing at least one species of metals of gallium, indium, lead, bismuth, aluminum, calcium and the like which are said to have an effect of increasing the hydrogen overvoltage and an effect of adjusting the particle shape during manufacturing of particles as the negative electrode active material, in the case of an electrolyte solution of the potassium hydroxide type, the amount of water in the battery is 0.31-0.57 mg (or 0.31-0.57 μ L at ordinary temperature) per a weight of 1 mg of mercuryless zinc, or in the case of an electrolyte solution of sodium hydroxide type, the amount of water in the battery is 0.32-0.59 mg (or 0.32-0.59 μ L at ordinary temperature) per a weight of 1 mg of mercuryless zinc, thereby it is possible to produce a coin or button type silver oxide battery in which the gas generation amount is not more than 0.03 μ L/g/day, and the self-discharge rate is not more than 4 %/year.

Further, in order to approximate the performance of the battery containing mercury, it is necessary to combine and use various techniques.

Mercuryless zinc alloy powder added with metals of indium, lead, bismuth, calcium, aluminum and the like and a gelling agent for holding moisture in the battery are used as a negative electrode combined agent, a collector having the outermost layer which is coated with zinc or metal having a hydrogen overvoltage higher than that of zinc is used, and an inhibitor selected from an indium compound, lead oxide, hydroxide of alkaline earth metal, and a surfactant having polyoxyethylene group is added into the electrolyte solution or the negative electrode active material, thereby it is possible to obtain a battery in which the hydrogen gas generation is less, and the electric characteristics are good.

Especially, with respect to the coating of the collector, an alloy layer containing zinc as an essential element and containing one or more species selected from indium, lead and tin as a selective element is provided, thereby it is possible to provide an alkaline battery which is advantageous from a viewpoint of cost in which the hydrogen gas generation can be suppressed.

Further, there are such problems due to elimination of mercury that the self-discharge increases and the capacity decreases, however, the zinc alloy layer of the collector is made thick to some extent, thereby it is possible to increase the capacity of the battery without changing the space in a battery can so much.

The indium compound such as indium sulfate, indium sulfamate, indium chloride and the like dissolves in a concentrated caustic alkali solution, which forms alkaline complex ion capable of cathode reduction referred to in the plating.

The alkaline complex ion of indium is subjected to zinc surface reduction representing a potential lower than a reduction potential of itself, and indium is immediately deposited as metal. In addition, the collector such as copper or the like contacts with zinc, so that it becomes to have the same potential as that of zinc, and indium is deposited in the same manner. When the surfaces of zinc and the collector are initially coated with indium, all of the surfaces become to have the potential of indium, and the electrochemical driving force is lost, so that further deposition of indium is ceased. However, when the zinc surface is newly

exposed by discharge, indium which exists as the alkaline complex ion is immediately reduced and deposited.

The inhibitor of the indium compound functions more effectively by adding the complexing agent. This is due to the fact that the alkaline complex ion of indium and hydrated indium ion are unstable, so that they precipitate with no complexing agent to be insoluble in the electrolyte solution, or even if they are dissolved, they precipitate as hydroxide due to minute environmental change, or they are apt to change into a viscous solution as polyion (those similar to the description in Inorganic chemistry Series 7, Coordinate Stereochemistry, written by Yoichi NIIMURA, published by Baihukan Co. Ltd., 65-66). In addition, the alkaline complex ion of indium and the hydrated indium ion are unstable and have low deposition potentials, but they have wide ranges of the potential, and the hydrogen gas generation is also induced during the deposition. Thus, when the complexing is performed with tartarate or EDTA, a stable complex ion is provided, it is possible to narrow the range of the deposition potential of indium, it is possible to separate from the deposition potential of hydrogen, and it is possible to deposit only indium without accompanying the hydrogen generation.

Other than indium, tin and lead also have smaller ionization tendency than zinc, so that when ion of these metals is allowed to exist in the electrolyte solution, it is possible to deposit these metals on the zinc surface. In addition, because the collector contacts with zinc, it becomes to have the same potential as that of zinc, and the above-mentioned metal is deposited. The function for further suppression of the corrosion and dissolution by mixing the indium compound, the tin compound containing tetravalent tin, lead oxide and the like is not certain, however, the following facts can be considered.

One is a fact that the alloy formation may take place when these metals are deposited on zinc and the collector. During the alloy formation, crystals of the deposited metal are made fine, and the surfaces of zinc and the collector are coated with homogeneous films having no deficiency. For example, in the case of an alloy of indium-tin, crystals are made fine then a composition near the eutectic point of about 50:50 in an atomic % is aimed. In the case of an alloy of the three component system, crystals become more complicated, and coarse formation of crystal grains is prevented, so that it becomes easy to obtain a homogeneous film.

The other is a fact that characteristics possessed by each of the metals can be simultaneously utilized by mixing. Especially, in the case of lead, if it is used alone, needle-like crystals are deposited, it is impossible to homogeneously coat the surfaces of zinc and the collector, and the effect is not so much. However, during the assembling of the battery, when the electrolyte solution containing the indium compound and the tin compound containing tetravalent tin is added previously, and subsequently the electrolyte solution containing lead oxide is added, then zinc and the collector are coated with indium and tin which form relatively homogeneous films, and then coated with a film having needle-like crystals of lead. The homogeneous films of indium and tin suppress the corrosion and dissolution of zinc or the collector, and the film of lead having needle-like crystals strengthens electric contact of zinc and the collector, and enhances shock resistance and discharge characteristics.

In the case of a negative electrode active material such as pure zinc and the like in which the hydrogen generation is large, it is difficult to form a homogeneous film due to the hydrogen generation which occurs as a competitive reaction with respect to the deposition of metal on the surface, and the effect of deposited metal becomes less. Thus, it is effective that such a negative electrode active material is used in which zinc is added with indium, bismuth, lead, aluminum, calcium, gallium and the like, and the hydrogen generation is suppressed to some extent.

Such an attempt has been most frequently made that zinc is added with metal having a high hydrogen overvoltage to form an alloy so as to suppress the corrosion and dissolution. It is known that the effect thereof is remarkably expressed especially when indium is added. When zinc is used in which indium is added in a relatively high concentration, for example, not less than 400 ppm, a part of indium and zinc is once dissolved due to the contact with copper or the like of the collector. It is considered that the corrosion and dissolution of zinc is suppressed by a mechanism that dissolved indium ion is then deposited on the collector, and a film of indium is formed on the collector.

However, in the initial stage of the contact between zinc and the collector, the amount of indium ion to be reduced on the collector is extremely small. Thus, when the one or more species of the compounds selected from the indium compound which is nobler than zinc, the tin compound containing tetravalent tin and lead oxide are added as the inhibitor into the electrolyte solution or the negative electrode active material, and the metal ion is allowed to exist in the electrolyte solution in an amount sufficient to coat zinc and the collector, then it is possible to immediately deposit lead onto zinc and the negative electrode collector, and it is possible to suppress the hydrogen generation.

When iron is abundant in zinc, there are provided many places in which iron is exposed on the zinc surface. Zinc and iron on the surface form a local battery in the electrolyte solution, zinc dissolves, and hydrogen is generated from iron. At the place in which hydrogen is generated, the coating film by the inhibitor is difficult to be formed, and it is difficult to obtain the effect of the inhibitor.

When mercuryless zinc powder in which the content of iron is not more than 4 ppm with respect to a weight of zinc is used, one or more species of the compounds selected from the indium compound nobler than zinc, the tin compound containing tetravalent tin and lead oxide are added as the inhibitor into the electrolyte solution or the negative electrode active material, and the ion is allowed to exist in the electrolyte solution in an amount sufficient to coat zinc and the collector, then it is possible to immediately deposit the film of metal nobler than zinc onto zinc and the negative electrode active material, and it is possible to suppress the corrosion and dissolution of zinc and the hydrogen generation accompanying therewith.

It is preferable that the adding amount of the inhibitor is about 10-1000 ppm with respect to the zinc powder. If the adding amount is less, it is impossible to sufficiently coat zinc and the collector, and there is such a bad effect that needle-like crystals penetrate through the separator to cause a short circuit.

It is generally said that gallium, indium, lead and bismuth have high hydrogen overvoltages, and when they are added to zinc, the hydrogen gas generation is suppressed. It is said that aluminum and calcium smoothen the surface during the production of zinc powder by atomization, reduce the surface area of the zinc powder, and suppress the hydrogen gas generation in the same manner.

In addition, although the function is not certain, the electrolyte solution is made as much as permitted by liquid leakage, thereby it is possible to suppress the hydrogen gas generation and the self-discharge. Further, by increasing the electrolyte solution, it also becomes possible to increase the absolute amount of the inhibitor to be added in order to suppress the hydrogen gas generation and the self-discharge, and it is considered that the inhibitor effect is also increased.

In order to allow the mercuryless battery to approximate the performance of a battery in which mercurated zinc is contained, it is necessary that features of various techniques are understood, and they are combined and used. Functions of individual techniques will be shown hereinafter.

Generally speaking, the addition of metals such as indium, lead, bismuth, calcium, aluminum and the like into zinc is to prevent the corrosion and dissolution and suppress the hydrogen gas generation. However, the role of each adding metal is varied. The metal having a high hydrogen overvoltage such as indium, lead and bismuth forms an alloy with zinc, increases the hydrogen overvoltage, and prevents the corrosion and dissolution. On the other hand, as a result of that calcium and aluminum smoothen the alloy surface during manufacturing by atomization, which level potential distribution, and decrease the surface area of zinc particles, so that they are effective in the prevention of the corrosion and dissolution. In addition, when hydrogen is generated on the zinc surface by the corrosion and dissolution, no inhibitor is supplied there and no effect of the inhibitor can be also exhibited. Also in this meaning, the alloy formation of zinc by the addition of metal of the present invention is important.

The cross-linked type polyacrylic water-absorbing polymer to be used as the gelling agent has a strong moisture holding property, which prevents the electrolyte solution from evaporation and unnecessary movement toward the separator and the positive electrode side. Thus, it is possible to allow the inhibitor to be distributed all over and suppress the increase in the internal resistance due to shortage of liquid at the last stage of discharge.

The reason why the layer of zinc or metal having a hydrogen overvoltage higher than that of zinc is formed at the outermost layer of the collector is to prevent the fact that the mercuryless zinc powder as the negative electrode active material contacts with copper of the collector to form a battery, and hydrogen gas is also generated from it.

In addition, when the layer of zinc or metal having a hydrogen overvoltage higher than that of zinc is formed after the press processing of a negative electrode can containing the collector, it is possible to shield impurities such as iron and the like adhered to the collector side during the processing.

Especially, it is effective to provide the alloy layer containing zinc as the essential element and containing one or more species selected from indium, lead and tin as the selective element on the collector surface. This is due to the fact that the potential of the collector surface becomes substantially the same as that of the mercuryless zinc surface as the negative electrode active material. Thus, even when the collector contacts with the mercuryless zinc powder as the negative electrode active material, it is avoided that a local battery is formed and hydrogen gas is generated from it.

It is supposed that, even if only zinc is plated on the collector, the potential difference may disappear and an effect may be expected, however, zinc deposited by plating or the like is extremely active and easy to be corroded, so that little effect is provided for suppressing the hydrogen gas formation. Thus, it is necessary to plate an alloy in which indium and lead are added into zinc by not less than several 10 ppm.

In the plating process, for the plating solution, the one in which several 10 to several 1000 ppm of the indium compound, lead compound and tin compound are added into general zinc plating solutions may be need, and for the positive electrode, an alloy of zinc may be used. For example, indium sulfate may be added in the case of a plating solution of the zinc sulfate type, while indium cyanide may be added in the case of a plating solution of the cyanide type.

As the inhibitor, there are the inorganic type and the organic type. Among them, the inorganic inhibitor is roughly classified into two species. One is a compound of metal which is nobler than zinc and has a higher hydrogen overvoltage. For example, it is the above-mentioned indium compound such as indium sulfate, indium sulfamate, indium chloride, indium hydroxide or the like, and they are dissolved in a concentrated caustic alkaline solution, and form alkaline complex ion capable of cathode reduction referred to in the plating. The alkaline complex ion of indium is reduced at the Zinc surface representing a potential lower than the reduction potential of itself, and indium is immediately deposited as metal. When zinc is initially coated with indium, all of the surface becomes to have the potential of indium, and the electrochemical driving force is lost, so that further deposition of indium is ceased. However, when the zinc surface is newly exposed by discharge, indium which exists as the alkaline complex ion is immediately reduced and deposited. Thus, there are provided such effects that the hydrogen gas generation is less even when the discharge is stopped on the way to store, and the self-discharge rate is made small.

Due to the fact that a potential lower than the reduction potential of itself is indicated also on the collector such as copper or the like contacting with zinc, the alkaline complex ion of indium is reduced, and indium is immediately deposited as for metal. However, the amount of the electrolyte solution is fairly less in an actual battery, so that the inhibitor is not homogeneously distributed on the collector to provide no metal coating, and when there are bubbles between the negative electrode combined agent and the collector, such a place only where no metal coating is provided at all is made, and it has been impossible to exhibit a sufficient effect. Lead monoxide and the tetravalent tin compound are included in the inorganic inhibitor of this type.

The other inorganic inhibitor is oxide and hydroxide of metal baser than zinc or non-metal. Although the function is not certain, effects are provided in the suppression of the hydrogen gas generation and the improvement in electric characteristics, and as the representative one, there is barium hydroxide or the like which is hydroxide of alkaline earth metal. However, when a collector which having no layer of zinc or metal having a hydrogen overvoltage higher than that of zinc for the outermost layer is used, the hydrogen gas generation due to the contact between the collector and zinc is too large, and little effect thereof is expressed.

The surfactant of organic type suppresses the corrosion and dissolution of zinc because the hydrophilic group adheres to the zinc surface and the hydrophobic group suppresses the approach of water and hydroxyl group to the surface. The effect is similar to that of a hydroxide of alkaline metal, and it is desirable to combine and use with the collector having the layer of zinc or metal having a hydrogen overvoltage higher than that of zinc for the outermost layer.

In addition, although the function is not certain, when polyoxyethylene alkylamide, barium hydroxide and the like are combined and used, a remarkable hydrogen gas generation suppressing effect is obtained. However, organic surfactants have a possibility to be excessively adhered onto the surfaces of zinc and the collector and inhibit the reduction on the surface in the combination with the indium compound or lead monoxide, so that it is desirable that the adding amount is made as small as possible within a range which can provide the effect.

It has been found out that there are roles of each one such that the coating of zinc or metal having a hydrogen overvoltage higher than that of zinc onto the collector suppresses the hydrogen gas generation due to the contact between zinc and the collector, the indium compound and lead monoxide suppress the self-discharge after partial discharge, and the alkaline earth hydroxide improves electric characteristics, and the maximum effect is obtained when they are combined and used. Namely, when the alkaline battery is manufactured by using the collector having the layer of zinc or metal having a hydrogen overvoltage higher than that of zinc for the outermost layer, it is possible to obtain the one in which the self-discharge rate is small before use and after partial discharge, and the electric characteristic is good.

When the collector is coated with a zinc alloy, the zinc alloy becomes the negative electrode active material as it is, so that by thickening the zinc alloy layer on the collector surface, it is possible to make up a battery capacity reduced by elimination of mercury. For example, when a collector having a diameter of 6 mm is plated by 10 μ m, the amount of the negative electrode active material increases by 2 mg because the specific gravity of zinc is 7.13. Provided that the amount of the negative electrode active material as zinc powder is 30 mg, it is possible to increase the capacity by about 6.7 % without changing the space in the battery can so much. However, as compared with the powdery negative electrode active material, the

surface area is small, so that no large current can be expected. However, it is most suitable for increasing the capacity of a batteries which perform minute discharge such as batteries for clocks and watches.

In addition, to thicken the zinc alloy plating is advantageous also in the processing of battery cans. Generally, the negative electrode can having the negative electrode collector is manufactured by punching a hoop material in many cases. In this case, if the zinc alloy has plated rather thickly beforehand at the side of hoop material for serving as the collector, the probability to expose metal such as for example copper as a base substrate during processing is reduced.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 shows the generated amount of hydrogen gas with respect to days passed.

Fig. 2 shows the hydrogen gas generated amount with respect to the indium sulfate concentration.

Fig. 3 shows current-potential curves at ordinary temperature in the potassium hydroxide solution.

Fig. 4 shows current-potential curves at ordinary temperature in the potassium hydroxide solution in which zinc oxide is dissolved up to approximate saturation.

Fig. 5 shows the hydrogen gas generation amount with respect to days passed.

Fig. 6 shows the hydrogen gas generation amount with respect to the indium sulfate concentration.

Fig. 7 is a figure showing the generation amount of hydrogen gas with respect to the amount of lead monoxide added into the electrolyte solution of the present invention.

Fig. 8 is a figure showing the generation amount of hydrogen gas with respect to the amount of lead monoxide added into the electrolyte solution.

Fig. 9 shows the hydrogen generation amount with respect to the amount of water added per 1 mg of zinc when the electrolyte solution of the potassium hydroxide type is used.

Fig. 10 shows the hydrogen generation amount with respect to the amount of water added per 1 mg of zinc when the electrolyte solution of the sodium hydroxide type is used.

Fig. 11 shows the generation amount of hydrogen gas with respect to days passed when the inhibitor of the present invention is added to the plated copper plate.

Fig. 12 shows the generation amount of hydrogen gas with respect to days passed when the inhibitor of the present invention is added to the non-plated copper plate.

Fig. 13 shows the change in the self-discharge rate with respect to the inhibitor concentration.

Fig. 14 shows the change in the self-discharge rate after 50 % depth discharge with respect to the inhibitor concentration.

Fig. 15 shows the generation amount of hydrogen gas with respect to days passed when the inhibitor of the present invention is added to the plated copper plate.

Fig. 16 shows the generation amount of hydrogen gas with respect to days passed when the inhibitor of the present invention is added to the non-plated copper plate.

Fig. 17 shows the change in the self-discharge rate with respect to the inhibitor concentration.

Fig. 18 shows the change in the self-discharge rate after 50 % depth discharge with respect to the inhibitor concentration.

Fig. 19 is a figure showing the hydrogen gas generation amount with respect to barium hydroxide.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention will be explained hereinafter in accordance with Embodiments.

The effect of the case in which the indium compound of the present invention is used will be explained in accordance with the Embodiments 1-3, and the Comparative example 1.

(Embodiment 1)

Into a specially prepared test tube having a volume of 25 ml and graduated to know the gas generation amount were added beforehand 2 g of zinc powder containing 500 ppm of each of bismuth, indium and lead manufactured by the atomization method, and a copper piece as the same material as a collector having an area of 0.6 cm² and a thickness of 0.1 mm, an electrolyte solution to be tested was added thereto to heat to 60 °C, and the volume of generated hydrogen gas was measured for 7 days. The number of test repeating was made to 10 times, and an average value thereof was used as a result. The electrolyte solution was prepared such that by making, a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide type, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate

saturation in the case of the sodium hydroxide type, as a base, an indium compound was added thereto.

Indium sulfate made by Nihon Kagaku Sangyo Co., Ltd., indium sulfamate 35 % solution also made by Nihon Kagaku Sangyo Co., Ltd., indium chloride made by Wako Pure Chemical Industries., Ltd., and indium cyanide made by Ito Pharmaceutical Co., Ltd. were used. The adding amount of the indium compound was 1000 ppm with respect to the electrolyte solution. Results are shown in the hydrogen generation amount in Table 1. The unit is μ L/g/day.

(Comparative example 1)

With the same test as in Embodiment 1, the measurements were performed for one in which no indium compound of the present invention was added, and one in which indium oxide was added by 1000 ppm with respect to the electrolyte solution. Indium oxide made by Kanto Chemical Co., Ltd. was used. Results are shown in the hydrogen generation amount in Table 1 in the same manner. The unit is μ L/g/day.

Table 1. Hydrogen evolution volume and comparison of discharge capacity of experimental cell

Examples of the invention	Electrolyte	Indium compound 1000ppm	Hydrogen evolution volume μ L/g/day	Comparison of discharge capacity
1	KOH	InSO ₄ Indium sulfate	24.29	102
2	KOH	In(NH ₂ SO ₃) ₃ Indium Sulfamate	30.51	102
3	KOH	Indium chloride	35.40	101
4	KOH	Indium cyanide	20.66	103
5	NaOH	InSO ₄ Indium sulfate	18.11	102
6	NaOH	In(NH ₂ SO ₃) ₃ Indium Sulfamate	27.43	102
7	NaOH	Indium chloride	29.67	102
8	NaOH	Indium cyanide	14.21	103
Examples for comparison				
9	KOH	-	516.63	100
10	KOH	Indium oxide	196.10	97
11	NaOH	-	420.36	100
12	NaOH	Indium oxide	150.88	97

In Fig. 1, the hydrogen gas generation amount with respect to the number of test days of the samples using the electrolyte solution of the potassium hydroxide series, is shown. It has been found that in the case of no addition of the indium compound shown in 1 in the figure, the hydrogen generation amount increases exponential functionally with respect to the number of test days. It is understood that in the case of addition of the indium sulfate in 2 in the figure which is the indium compound of the present invention, the generation of hydrogen gas is suppressed. In the case of addition of the indium oxide in 3 in the figure, the hydrogen gas generation in the former half is much. This is considered to be due to the fact that the solubility of indium oxide is bad, the supply of indium ion to the electrolyte solution is not sufficient, and a long time is required to coat the surfaces of zinc and copper.

In addition, it is considered that the reason why a bad result is given in the former half as compared with no addition one is that indium oxide also contacts with zinc and the collector, and a local battery is formed. Indium cyanide presented a good result from a viewpoint of characteristics, however, it is desirable not to use it because there is possibility of occurrence of a new environmental pollution problem.

(Embodiment 2)

Into a positive electrode can were added a part of an electrolyte solution and 116 mg of a pellet (silver oxide content of 98 %) which was molded by adding a combined agent into silver oxide, and a separator of polyethylene and a separator of cellophane were set place. Next, a gasket of nylon was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, 30 mg of zinc powder, an inhibitor and the like were added, the remainder of the electrolyte solution was added dropwise, and then a negative electrode can was set place to perform sealing to manufacture 100 pieces for every kind of button type silver oxide batteries. The same zinc as in Embodiment 1 and the electrolyte solution in Table 1 were used, and the battery size was made as the SR621 type.

However, the adding amount of indium was 1000 ppm with respect to the amount of zinc. Results are shown in the discharge index in Table 1. Discharge characteristics were measured using a resistance of 200 Ω by means of the direct current method when the electrolyte solution was the potassium hydroxide series or the pulse method when it was the sodium hydroxide. In any case, no addition of the inhibitor was regarded to have a discharge index of 100. As understood also from the results, it has been found that the indium compound of the present invention is effective also in the discharge characteristics.

(Embodiment 3)

Using a gasket of polypropylene specially prepared for allowing hydrogen to penetrate in the same manner as Embodiment 2, a button type silver oxide battery was manufactured. Ten pieces of the manufactured silver oxide batteries were put into a vessel made of glass filled with liquid paraffin in a high temperature tank, a collecting tube having graduations at an upper portion was attached, and the amount of generated hydrogen gas was measured. This state was maintained at 60 °C for 20 days said which as to correspond to a term of about 1 year, and the hydrogen gas generation amount after 20 days was investigated. With respect to the inhibitor, the test was performed using indium sulfate at a concentration of 10 ppm to 5 % with respect to the weight of zinc. Evaluation results are shown in Fig. 2. According to the figure, it is understood that the inhibitor effectively works at 100 ppm to 1 %.

The gasket of polypropylene was returned to one made of nylon, and a button type silver oxide battery was manufactured in a range of an indium sulfate concentration of 100 ppm to 1 % in which the gas generation was less. Ten pieces of the manufactured batteries were put into a vessel made of glass filled with liquid paraffin in a high temperature tank maintained at 25 °C in the same manner, and a collecting tube for generating hydrogen gas was attached to the upper portion. Hydrogen gas generation, expansion of the can and liquid leakage were not observed after 20 days at 60 °C. For indium sulfate, it was effective in the concentration of 100 ppm to 1 % with respect to zinc, however, the same effect was observed also in the case of other indium compounds by adjusting the mole number and determining the concentration range.

In the actual trial production of batteries, there is given a value smaller than the hydrogen gas generation amount in the experiment in Embodiment 1. In respect to this fact, it is considered that the copper piece used in Embodiment 1 is different from a structure of an actual collector, and a part of generated hydrogen gas has been consumed by the reduction of silver oxide and the like. The method in Embodiment 1 is different in the amount order of hydrogen gas generation, but it is sufficient as a method for observing substituted characteristics for predicting hydrogen generation in an actual battery.

In accordance with Embodiments 4-7, the effect of the case in which the indium compound and the complexing agent of the present invention are used will be explained.

(Embodiment 4)

For an electrolyte solution of 30 % by weight of potassium hydroxide, using a platinum electrode of 1 cm² as the working electrode, a platinum electrode of 1 cm² as the counter electrode in the same manner, and a comparative electrode HC-205C made by Toa Electronics Ltd. as the reference electrode, the current-potential curve during deposition of indium was determined. The potential was scanned from the plus side to the minus side at a speed of 100 mV/second. The current for each potential represents the

reaction at the electrode, and for example, if the reduction reaction such as deposition of indium and generation of hydrogen gas occurs, a current caused thereby flows.

Results are shown in Fig. 3. 1 in the figure is a case where indium sulfamate was added to the electrolyte solution by 0.1 mol/L, in this case, the current began to flow from the vicinity of -1.1 V with respect to the reference electrode, and indium was deposited accompanying hydrogen generation. 2 is a current-potential curve of a case in which 0.1 mol/L of potassium tartarate was added to the electrolyte solution, and then 0.1 mol/L of indium sulfamate was added. It was found that indium was deposited at the vicinity of -1.4 V, and hydrogen was generated under the vicinity of -2.0 V. This is considered such that since indium having a high hydrogen overvoltage was deposited on the electrode surface, the potential for hydrogen generation was shifted toward the minus side.

Thereby, the potential of deposition of indium could be separated from that of hydrogen generation. It is considered that also in the case of an actual battery, if indium is deposited previously, the hydrogen gas generation further becomes smaller. In the case where indium sulfamate and potassium tartarate are added to the electrolyte solution, when indium sulfamate is added previously, indium hydroxide and polyion are generated to become cloudy, while when potassium tartarate is added previously, no turbidity occurs.

In 3 in Fig. 3 is shown a current-potential curve of a case where indium sulfamate was added to the electrolyte solution by 0.1 mol/L, and then 0.1 mol/L of potassium tartarate was added. A small peak probably considered as indium is observed in the vicinity of -1.4 V in the figure, however, indium is deposited accompanying hydrogen generation from the vicinity of -1.1 V with respect to the reference electrode in the same manner as the case of no addition of potassium tartarate, and the effect of addition of the complexing agent becomes small. This also can be said for other indium compounds and complexing agents, and in order to immediately form the complex ion of indium and prevent precipitation, it is necessary to dissolve the complexing agent previously in the electrolyte solution.

25 (Embodiment 5)

For an electrolyte solution of 30 % by weight of potassium hydroxide in which zinc oxide was added up to approximate saturation, using a platinum electrode of 1 cm² as the working electrode, a platinum electrode of 1 cm² as the counter electrode in the same manner, and a comparative electrode HC-205C made by Toa Electronics Ltd. as the reference electrode, the current-potential curve during deposition of indium was determined. Results are shown in Fig. 4. 4 in the figure is a case where indium sulfamate was added to the electrolyte solution by 0.1 mol/L, and metal was deposited accompanying hydrogen generation from the vicinity of -1.56 V with respect to the reference electrode. The deposition potential of metal is lower than that of a case of no addition of zinc oxide, so that there is a possibility that it is an alloy of zinc and indium.

5 is a current-potential curve of a case where 0.1 mol/L of potassium tartarate was added to the electrolyte solution, and then 0.1 mol/L of indium sulfamate was added. There is a peak considered as the deposition of indium at the vicinity of -1.5 V, and then metal was deposited accompanying hydrogen generation from the vicinity of -1.56 V. In order to confirm the deposited substance at the vicinity of -1.5 V, the potential was held at -1.5 V for 30 seconds, and the deposited metal was immersed in nitric acid. If it was zinc, it might be instantly dissolved, but it was difficult to dissolve, so that the deposited substance is postulated to be indium or an alloy having a large amount of indium.

As a result of measurement, the electrode potential of zinc was -1.509 V with respect to the reference electrode. It just corresponds to a potential at which only indium or an alloy having a large amount of indium is deposited in a solution containing a complexing agent. Namely, indium or the alloy having a large amount of indium is deposited on zinc and the collector contacting with zinc, which prevents zinc and the collector from corrosion and dissolution.

(Embodiment 6)

50 Into a specially prepared test tube having a volume of 25 ml and graduated to know the gas generation amount were added beforehand 2 g. of zinc powder containing 500 ppm of each of bismuth, indium and lead manufactured by the atomization method, and a copper piece as the same material as the collector having an area of 0.6 cm² and a thickness of 0.1 mm, an electrolyte solution to be tested was added thereto to heat to 60 °C, and the volume of generated hydrogen gas was measured for 7 days. The number of repetition of the test was made 10, and an average value thereof was used as a result.

The electrolyte solution was prepared such that by making a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium

hydroxide series, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide series, as a base, and a complexing agent and an indium compound were added thereto.

Indium sulfate made by Nihon Kagaku Sangyo Co., Ltd., indium sulfamate 35 % solution also made by Nihon Kagaku Sangyo Co., Ltd., indium chloride made by Wako Pure Chemical Industries., Ltd., and indium cyanide made by Ito Pharmaceutical Co., Ltd. were used

The adding amount of the indium compound was 1000 ppm with respect to the electrolyte solution, and the adding amount of the complexing agent was made 2-fold excessive in a mole ratio of indium ion.

Results in the case where the indium compound was added without adding of the complexing agent are shown in Table 2. The hydrogen gas generation amount is shown in Table 3 in the case where the complexing agent was added. The hydrogen generation index is the value in which the hydrogen generation amount in the test using the same electrolyte solution except for the complexing agent is regarded as 100.

When the hydrogen gas generation amount is observed, so as postulated from the current-potential curve, it is understood that the hydrogen generation is suppressed. Although the effect is observed also in the cyanide series, however, the cyanide type has a possibility to lead a new environmental pollution problem, so that it is safe not using this.

Table 2

Hydrogen evolution volume without complexing agent and comparison of discharge capacity of experimental cell					
Examples of the invention	Electrolyte	Indium compound 1000ppm	Complexing agent	Hydrogen evolution volume μ L/g/day	Comparison of discharge capacity
1	KOH	Indium sulfate	-	24.29	100
2	KOH	Indium sulfamate	-	30.51	100
3	KOH	Indium chloride	-	35.40	100
4	KOH	Indium cyanide	-	20.66	100
5	NaOH	Indium sulfate	-	18.11	100
6	NaOH	Indium sulfamate	-	27.43	100
7	NaOH	Indium chloride	-	29.67	100
8	NaOH	Indium cyanide	-	14.21	100

Table 3. Comparison of hydrogen evolution volume with complexing agent and comparison of discharge capacity of experimental cell

Examples of the invention	Electrolyte	Indium compound 1000ppm	Complexing agent	Hydrogen evolution volume $\mu\text{L/g/day}$	Comparison of discharge capacity
9	KOH	Indium sulfate	Potassium tartrate	98	102
10	KOH	Indium sulfamate	Potassium tartrate	99	101
11	KOH	Indium chloride	Potassium tartrate	98	103
12	KOH	Indium cyanide	Potassium tartrate	99	102
13	NaOH	Indium sulfate	Sodium tartrate	98	102
14	NaOH	Indium sulfamate	Sodium tartrate	97	103
15	NaOH	Indium chloride	Sodium tartrate	98	101
16	NaOH	Indium cyanide	Sodium tartrate	98	102
17	KOH	Indium sulfate	EDTA	98	102
18	KOH	Indium sulfamate	EDTA	98	102
19	KOH	Indium chloride	EDTA	99	103
20	KOH	Indium cyanide	EDTA	98	103
21	NaOH	Indium sulfate	EDTA	97	101
22	NaOH	Indium sulfamate	EDTA	99	102
23	NaOH	Indium chloride	EDTA	98	102
24	NaOH	Indium cyanide	EDTA	97	102
25	KOH	Indium sulfate	Glycine	99	101
26	KOH	Indium sulfamate	Glycine	99	100
27	KOH	Indium chloride	Glycine	99	101
28	KOH	Indium cyanide	Glycine	99	101
29	NaOH	Indium sulfate	Glycine	99	101
30	NaOH	Indium sulfamate	Glycine	99	100
31	NaOH	Indium chloride	Glycine	99	101
32	NaOH	Indium cyanide	Glycine	98	101
33	KOH	Indium sulfate	Potassium cyanide	97	103
34	KOH	Indium sulfamate	Potassium cyanide	97	103
35	KOH	Indium chloride	Potassium cyanide	97	103
36	KOH	Indium cyanide	Potassium cyanide	97	104
37	NaOH	Indium sulfate	Sodium cyanide	97	103
38	NaOH	Indium sulfamate	Sodium cyanide	97	102
39	NaOH	Indium chloride	Sodium cyanide	97	102
40	NaOH	Indium cyanide	Sodium cyanide	96	103

(Embodiment 7)

Into a positive electrode can were added a part of an electrolyte solution and 116 mg of a pellet (silver oxide content of 98 %) which was molded by putting a combined agent into silver oxide, and a separator of polyethylene and a separator of cellophane were set place. Next, a gasket of nylon was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, 30 mg of zinc powder, an inhibitor and the like were added, the remainder of the electrolyte solution was added dropwise, and then a negative electrode can was set place to perform sealing to manufacture 100 pieces for every kind of button type silver oxide batteries. The same zinc as in Embodiment 6 and the electrolyte solution in Table 2 were used, and the battery size was made as the SR621 type.

However, the adding amount of indium was made 1000 ppm with respect to the amount of zinc, and the adding amount of the complexing agent was made 2-fold in a mole ratio of indium ion. Discharge characteristics were measured using a resistance of 200 Ω by means of the direct current method when the electrolyte solution was the potassium hydroxide series or the pulse method when it was the sodium hydroxide series. In any case, no addition of the complexing agent was regarded to have a discharge index of 100. Results are shown with the right discharge index in Table 2. The effect of glycine is rather small, however, the effect was observed for all of the tested complexing agents.

Explanation has been made on the basis of Embodiment for the representative indium compound and the complexing agent as described above, however, the effects of the indium compound and the complexing agent can be easily postulated if the current-potential curve is determined in the same manner as shown in Embodiments 4 and 5. It is needless to say that other indium compounds and complexing

agents exhibiting the same effects can be applied to the present invention. In addition, an example for the silver battery has been described in Embodiment 4, however, the same effect can be expected even in the case of alkaline manganese batteries using zinc, air batteries and the like.

In addition, in the case of a negative electrode active material such as pure zinc and the like in which hydrogen generation is large, a homogeneous film is difficult to be formed due to hydrogen generation which occurs on the surface as a competitive reaction against indium deposition, and the effect of indium deposition becomes small. Thus, it is effective to use a negative electrode active material in which the hydrogen generation is suppressed to some extent by adding indium, bismuth, lead, aluminum, gallium, calcium and the like into zinc.

In Embodiment 8-11, the effect in the case in which two or more species selected from the indium compound, the tin compound containing tetravalent tin and lead oxide are used as the inhibitor of the present invention will be explained. (Embodiment 8)

Into a specially prepared test tube having a volume of 25 ml graduated to know the gas generation amount were added beforehand 2 g of zinc powder containing 500 ppm of each of bismuth, indium and lead manufactured by the atomization method, and a copper piece as the same material as a collector having an area of 0.6 cm² and a thickness of 0.1 mm, an electrolyte solution to be tested was added thereto to heat to 60 °C, and the volume of generated hydrogen gas was measured for 7 days.

The number of repetition of the test was made 10, and the average value thereof was used as a result. The electrolyte solution was prepared such that by adding a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide series, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide series, as a base, and compounds selected from indium sulfate, indium sulfamate, sodium stannate and lead oxide were added thereto.

Indium sulfate and sodium stannate made by Nihon Kagaku Sangyo Co., Ltd., indium sulfamate 35 % solution also made by Nihon Kagaku Sangyo Co., Ltd., and lead oxide made by Wako Pure Chemical, Ltd. were used. The adding amount of the compounds was made 1000 ppm in total with respect to the electrolyte solution. Results are shown in the hydrogen generation amount in Table 4.

Table 4. Hydrogen evolution volume and comparison
of discharge capacity of experimental cell

Examples of the invention	Amount of added compound				Electro- lyte	The number of copper pieces soaked in electrolyte	Hydrogen evolution μ /g/day	Comparison of discharge capacity
	Indium sulfate	Indium sulfamate	Lead oxide	Sodium stannet				
1	1000				KOH	0	37.86	—
2		1000			KOH	0	27.14	—
3			1000		KOH	0	157.36	—
4				1000	KOH	0	26.81	—
5	1000				KOH	5	24.29	102
6		1000			KOH	5	30.51	102
7			1000		KOH	5	55.56	103
8				1000	KOH	5	39.15	101
9	500		500		KOH	5	21.43	103
10	500			500	KOH	5	35.71	102
11			500	500	KOH	5	50.00	103
12		500	500		KOH	5	18.57	103
13		500		500	KOH	5	46.43	102
14	333		333	333	KOH	5	42.86	103
15		333	333	333	KOH	5	42.86	103
16	1000				NaOH	0	35.26	—
17		1000			NaOH	0	23.85	—
18			1000		NaOH	0	103.45	—
19				1000	NaOH	0	24.17	—
20	1000				NaOH	5	22.14	102
21		1000			NaOH	5	27.51	102
22			1000		NaOH	5	48.23	103
23				1000	NaOH	5	35.12	101
24	500		500		NaOH	5	18.82	103
25	500			500	NaOH	5	31.48	102
26			500	500	NaOH	5	44.22	103
27		500	500		NaOH	5	16.11	103
28		500		500	NaOH	5	39.64	102
29	333		333	333	NaOH	5	37.26	103
30		333	333	333	NaOH	5	35.12	103
31					KOH	5	516.63	/100

50

For comparison, experiment results of no addition of the copper plate in the same experiment as Embodiment 8 are shown in 1-4 and 16-19 in Table 4. In addition, a result of no addition of the compound is shown in 31 in Table 4. The hydrogen gas generation amount with respect to the number of test days for 31 in Table 4 and 12 in Table 4 as a combination of indium sulfamate and lead oxide whose gas generation was small is shown in Fig. 5. It is understood that in the case of no addition of the compound, the hydrogen generation amount increases with respect to the number of test days exponential functionally, and it is understood that in the case of the addition of indium sulfamate and lead oxide, the generation of hydrogen gas is suppressed.

As seen from the data of 3 and 4 in Table 4, lead oxide provides a relatively large amount of hydrogen generation. This is considered to be due to the fact that when lead is deposited, needle-like crystals are generated to become impossible to homogeneously coat the surfaces of zinc and the collector. From a viewpoint of the combination, it is understood that the combination of the compound of indium and lead oxide is especially good, which suppress the hydrogen generation, and the discharge characteristics are improved.

(Embodiment 9)

Into a positive electrode can were added a part of an electrolyte solution and 116 mg of a pellet (silver oxide content of 98 %) which was molded by adding a combined agent into silver oxide, and a separator of polyethylene and a separator of cellophane were set place. Next, a gasket of nylon was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, and 30 mg of zinc powder were added, the remainder of the electrolyte solution added with the compound selected from indium sulfate, indium sulfamate, sodium stannate and lead oxide was added dropwise, and then a negative electrode can was set place to perform sealing to manufacture 100 pieces for every kind of button type silver oxide batteries.

The same zinc as in Embodiment 8 and the electrolyte solution composition in Table 4 were used, and the battery size was made as the SR621 type. However, the adding amount of the compound was 1000 ppm with respect to the zinc amount. Results are shown in the discharge index in Table 4. Discharge characteristics were measured using a resistance of 200 Ω by means of the direct current method when the electrolyte solution was the potassium hydroxide series or the pulse method when it was the sodium hydroxide series. In any case, no addition of the inhibitor was regarded to have a discharge index of 100. As understood also from the results, it has been found that the present invention is effective also in the discharge characteristics.

Especially, one in which lead oxide is added has a discharge index superior to those of others owing to the improvement in the electric contact by needle-like crystals. From a viewpoint of the combination, it is understood that the combination of the compound of indium and lead oxide is especially good, which suppress the hydrogen generation, and the discharge characteristics are improved. It can be supposed that indium suppresses the gas generation from zinc and the collector, and lead improves the discharge characteristics.

(Embodiment 10)

Using a gasket of polypropylene specially prepared for allowing hydrogen to penetrate in the same manner as Embodiment 9, a button type silver oxide battery was manufactured. Ten pieces of the manufactured silver oxide batteries were put into a vessel made of glass filled with liquid paraffin in a high temperature tank, a collecting tube having graduations at an upper portion was attached, and the amount of generated hydrogen gas was measured. In this state, it was maintained at 60 °C for 20 days, which is so as said to correspond to a term of about 1 year, and the hydrogen gas generation amount after 20 days was investigated. With respect to the adding compound, the test was performed using indium sulfate and lead oxide in a weight ratio of 1:1 at a concentration of 10 ppm to 5 % in total with respect to the weight of zinc. Evaluation results are shown in Fig. 6. According to the figure, it is understood that the adding compound effectively works at 50 ppm to 1 %.

The gasket of polypropylene was returned to one made of nylon, and a button type silver oxide battery was manufactured in a range of concentration of 50 ppm to 1 % using indium sulfate and lead oxide in which the gas generation was less. Ten pieces of the manufactured batteries were put in a vessel made of glass filled with liquid paraffin in a high temperature tank maintained at 60 °C in the same manner, and a collecting tube for generating hydrogen gas was attached to the upper portion. Hydrogen gas generation, expansion of the can and liquid leakage were not observed after 20 days at 60 °C.

Indium sulfate and lead oxide were effective for zinc at the concentration of 50 ppm to 1 %, however, the same effect was observed also in the case of other combinations of compounds by adjusting the mole number and determining the concentration range.

In the actual trial production of batteries, there is given a value smaller than the hydrogen gas generation amount in the experiment in Embodiment 8. That is, it is considered that the copper piece used in Embodiment 8 is different from a structure of an actual collector, a part of generated hydrogen gas has been consumed by the reduction of silver oxide, and the like. The method in Embodiment 8 is different in the amount order of hydrogen gas generation, but it is sufficient as a method for observing substituted

characteristics for predicting hydrogen generation in an actual battery.

(Embodiment 11)

5 Using a gasket of polypropylene specially prepared for allowing hydrogen to penetrate in the same manner as Embodiment 9, a button type silver oxide battery was manufactured. However, two kinds of electrolyte solutions indium sulfate and lead oxide of 50 ppm with respect to zinc, were prepared, and they were added by dividing into two times with changing the order. With respect to the manufactured button type silver oxide batteries, the test was performed at 60 °C for 20 days in the same manner as
10 Embodiment 9. The hydrogen generation amount was 0.10 μ L/g/day for one in which indium sulfate was added previously, and it was 0.15 μ L/g/day for one in which lead oxide was added previously. As understood also from the results, one in which indium sulfate was added previously has a larger effect for the hydrogen gas generation.

15 In accordance with Embodiments 12-14, the effect of the case where zinc containing indium is used with one or more species of compounds selected from the indium compound, the tin compound containing tetravalent tin and lead oxide as the inhibitor of the present invention will be explained.

(Embodiment 12)

20 Into a specially prepared test tube having a volume of 25 ml graduated to know the gas generation amount were put beforehand 2 g of zinc powder containing indium manufactured by the atomization method, and a copper piece as the same material as a collector having an area of 0.6 cm² and a thickness of 0.1 mm, an electrolyte solution to be tested was added thereto to heat to 60 °C, and the volume of generated hydrogen gas was measured for 7 days. The number of repetition of the test was 10, and an
25 average value thereof was used as a result. The electrolyte solution was prepared such that by making a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide series, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide series as a base, and the inhibitor of the present invention such as lead monoxide and the like was added
30 thereto.

In Table 5, the adding amount of indium into the zinc powder and the generation amount of hydrogen gas with respect to the adding amount of lead monoxide are shown. The adding amount of lead monoxide is shown by ppm with respect to the electrolyte solution. Results are also shown in the hydrogen generation amount in Table 5.

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Table 5. Hydrogen volume generated from zinc powder

Experimental No.	Electrolyte	Indium content of zinc powder ppm	Concentration of PbO in electrolyte ppm	Hydrogen evolution volume $\mu\text{L/g/day}$
1	KOH	200	0	603.31
2	KOH	200	10	398.88
3	KOH	200	50	102.56
4	KOH	200	100	14.29
5	KOH	200	500	87.01
6	KOH	200	1000	105.70
7	KOH	200	5000	122.43
8	KOH	500	0	516.92
9	KOH	500	10	30.01
10	KOH	500	50	20.13
11	KOH	500	100	6.21
12	KOH	500	500	14.73
13	KOH	500	1000	40.56
14	KOH	500	5000	143.32
15	KOH	1800	0	99.21
16	KOH	1800	10	10.05
17	KOH	1800	50	5.33
18	KOH	1800	100	5.05
19	KOH	1800	500	10.11
20	KOH	1800	1000	31.01
21	KOH	1800	5000	142.58
22	NaOH	200	0	55.01
23	NaOH	200	10	327.38
24	NaOH	200	50	70.25
25	NaOH	200	100	10.67
26	NaOH	200	500	62.98
27	NaOH	200	1000	85.22
28	NaOH	200	5000	110.12
29	NaOH	500	0	501.39
30	NaOH	500	10	25.52
31	NaOH	500	50	16.33
32	NaOH	500	100	3.99
33	NaOH	500	500	9.63
34	NaOH	500	1000	25.42
35	NaOH	500	5000	103.44
36	NaOH	1800	0	82.54
37	NaOH	1800	10	10.11
38	NaOH	1800	50	4.87
39	NaOH	1800	100	4.51
40	NaOH	1800	500	8.73
41	NaOH	1800	1000	24.42
42	NaOH	1800	5000	121.12

The hydrogen gas generation results in the electrolyte solution of the potassium hydroxide series are shown in Fig. 7. 1 in the figure is a case where zinc powder added with 200 ppm of indium was used, and there is an effect of suppressing hydrogen generation at an adding amount of lead monoxide of about 100-400 ppm. 2 is a case where zinc powder added with 500 ppm of indium was used, and there is an effect of suppressing hydrogen generation at an adding amount of lead monoxide of not less than 10 ppm. 3 is a case where zinc powder added with 1800 ppm of indium was used, and there is an effect in a concentration range equivalent to that in 2. Especially when indium is added in a high concentration in such a manner, the effect at an adding amount of lead monoxide of not more than 100 ppm is remarkable, and it is possible to reduce the adding amount of lead as an environmental pollution substance.

(Embodiment 13)

Into a positive electrode can were added a part of an electrolyte solution and 116 mg of a pellet (silver oxide content of 98 %) which was molded by adding a combined agent into silver oxide, and a separator of polyethylene and a separator of cellophane were set place. Next, a gasket of nylon was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, 30 mg of zinc powder, an inhibitor and the like were added, the remainder of the electrolyte solution was added dropwise, and then a negative electrode can was set place to perform sealing to manufacture 100 pieces for every kind of button type silver oxide batteries. The battery size was made as SR621 type. The electrolyte solution was made again so as to allow the adding amount of lead monoxide to be an amount with respect to the zinc amount. They were stored at 60 °C for 20 days, and results of measurement of the self-discharge rate are shown in Table 6.

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Table 6. Self-discharge rate of cell and hydrogen evolution rate

Experimental No.	Electro-lyte	Indium content of zinc powder ppm	Amount of PbO added to electrolyte (PbO weight to zinc weight) ppm	Self-discharge rate %	Hydrogen evolution rate $\mu\text{L/g/day}$
1	KOH	200	0	12.1	1.23
2	KOH	200	10	9.9	0.95
3	KOH	200	50	7.4	0.88
4	KOH	200	100	4.1	0.50
5	KOH	200	500	8.7	0.71
6	KOH	200	1000	9.1	0.91
7	KOH	200	5000	15.8	1.56
8	KOH	500	0	11.2	1.11
9	KOH	500	10	4.2	0.56
10	KOH	500	50	3.5	0.41
11	KOH	500	100	2.5	0.35
12	KOH	500	500	2.9	0.32
13	KOH	500	1000	3.6	0.47
14	KOH	500	5000	18.9	1.49
15	KOH	1800	0	10.3	1.02
16	KOH	1800	10	2.2	0.30
17	KOH	1800	50	1.7	0.25
18	KOH	1800	100	1.5	0.22
19	KOH	1800	500	2.1	0.28
20	KOH	1800	1000	3.7	0.39
21	KOH	1800	5000	16.6	1.70
22	NaOH	200	0	10.3	1.15
23	NaOH	200	10	8.2	0.90
24	NaOH	200	50	6.5	0.74
25	NaOH	200	100	3.3	0.42
26	NaOH	200	500	7.5	0.72
27	NaOH	200	1000	8.1	0.86
28	NaOH	200	5000	13.2	1.35
29	NaOH	500	0	10.2	1.00
30	NaOH	500	10	4.0	0.44
31	NaOH	500	50	2.9	0.32
32	NaOH	500	100	2.2	0.30
33	NaOH	500	500	2.3	0.29
34	NaOH	500	1000	3.1	0.48
35	NaOH	500	5000	16.7	1.38
36	NaOH	1800	0	8.9	0.90
37	NaOH	1800	10	1.9	0.31
38	NaOH	1800	50	1.3	0.21
39	NaOH	1800	100	0.9	0.19
40	NaOH	1800	500	1.5	0.25
41	NaOH	1800	1000	2.9	0.33
42	NaOH	1800	5000	12.3	1.54

As seen also from the results, it is understood that the reduction in the battery capacity is suppressed by adding 100-1000 ppm of lead monoxide in the case of the zinc powder added with 500 ppm of indium, or 10-1000 ppm of lead monoxide in the case of the zinc powder added with 1800 ppm of indium. The decrease in the battery capacity by the addition of lead monoxide at the high concentration is considered to be due to the fact that needle-like crystals of lead grew on account of excessive addition, and a short circuit

might be caused.

(Embodiment 14)

5 A button type silver oxide battery was manufactured in the same manner as Embodiment 13 except that a gasket of polypropylene specially prepared for allowing hydrogen to penetrate was used. Ten pieces of the manufactured silver oxide batteries were put into a vessel made of glass filled with liquid paraffin in a high temperature tank, a collecting tube having graduations at an upper portion was attached, and the amount of generated hydrogen gas was measured. In this state, it was maintained at 60 °C for 20 days, which is said to correspond to a term of about 1 year, and the hydrogen gas generation amount after 20 days was investigated. Results are shown in Table 6.

As understood from Embodiments, the hydrogen generation is mild and the self-discharge is also less in the case of the sodium hydroxide series as compared with the potassium hydroxide series with respect to the electrolyte solution.

15 In addition, the more the indium content in the zinc powder is, the better results are obtained, and the smaller amount the adding amount of lead monoxide is.

In the actual trial production of batteries, there is given a value smaller than the hydrogen gas generation amount in the experiment in Embodiment 12. In this fact, it is considered that the copper piece used in Embodiment 12 is different from a structure of an actual collector, a part of generated hydrogen gas 20 has been consumed by the reduction of silver oxide and the like.

The method in Embodiment 12 is different in the amount order of hydrogen gas generation, but it is sufficient as a method for observing substituted characteristics for predicting hydrogen generation in an actual battery.

The same experiment was performed for indium compounds, tin compounds containing tetravalent tin, and mixtures thereof other than lead monoxide. Results thereof are substantially the same as those in the case of lead monoxide, and it has been found that the reduction in the battery capacity is suppressed by adding 100-1000 ppm of the inhibitor in the case of the zinc powder added with 500 ppm of indium, or 10-1000 ppm of the inhibitor in the case of the zinc powder added with 1800 ppm of indium.

In accordance with Embodiments 15-17, the effect will be explained in the case where zinc containing 30 not more than 4 ppm of iron and one or more species of compounds selected from the indium compound, the tin compound containing tetravalent tin, and lead oxide as the inhibitor of the present invention were used will be explained.

(Embodiment 15)

35 As for zinc powder, one containing 100 ppm of gallium manufactured by the atomization method, 200 ppm of indium, 500 ppm of lead and 450 ppm of aluminum, and further containing 5 ppm of iron was used. The concentration of iron in the zinc powder was adjusted by performing iron removal by means of a magnet. The concentration after the iron removal was confirmed by the atomic absorption method.

40 The hydrogen generation test was performed such that into a specially prepared test tube having a volume of 25 ml and graduated to know the gas generation amount were put beforehand 2 g of zinc powder and a copper piece as the same material as a collector having an area of 0.6 cm² and a thickness of 0.1 mm, an electrolyte solution to be tested was added thereto to heat to 60 °C, and the volume of generated hydrogen gas was measured for 7 days. The number of repetition of the test was 10, and an average value 45 thereof was used as a result. The electrolyte solution was prepared such that by making a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide series, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide series, as a base lead monoxide or the like was added thereto as the inhibitor.

50 The concentration of iron in the zinc powder and the generation amount of hydrogen gas with respect to the adding amount of the inhibitor are shown in Table 7. The adding amount of the inhibitor is shown by ppm with respect to the electrolyte solution. Results are also shown in the hydrogen generation amount in Table 7.

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Table 7. Hydrogen volume generated from zinc powder

Experimental No.	Electro- lyte	Content of Fe in zinc powder ppm	Concentration of PbO in electrolyte ppm	Hydrogen evolution rate $\mu\text{L/g/day}$
1	KOH	5	0	570.00
2	KOH	5	10	53.75
3	KOH	5	50	56.25
4	KOH	5	100	79.29
5	KOH	5	500	120.85
6	KOH	5	1000	158.92
7	KOH	5	5000	284.28
8	KOH	4	0	508.94
9	KOH	4	10	21.14
10	KOH	4	50	16.13
11	KOH	4	100	20.15
12	KOH	4	500	45.21
13	KOH	4	1000	50.56
14	KOH	4	5000	201.73
15	KOH	2	0	516.92
16	KOH	2	10	12.50
17	KOH	2	50	12.50
18	KOH	2	100	14.29
19	KOH	2	500	30.21
20	KOH	2	1000	53.57
21	KOH	2	5000	162.85
22	NaOH	5	0	423.85
23	NaOH	5	10	42.36
24	NaOH	5	50	40.23
25	NaOH	5	100	66.34
26	NaOH	5	500	100.25
27	NaOH	5	1000	132.58
28	NaOH	5	5000	278.90
29	NaOH	4	0	396.37
30	NaOH	4	10	19.36
31	NaOH	4	50	12.72
32	NaOH	4	100	15.64
33	NaOH	4	500	38.38
34	NaOH	4	1000	47.25
35	NaOH	4	5000	190.01
36	NaOH	2	0	43.56
37	NaOH	2	10	11.23
38	NaOH	2	50	10.12
39	NaOH	2	100	12.93
40	NaOH	2	500	27.81
41	NaOH	2	1000	42.21
42	NaOH	2	5000	130.10

Hydrogen gas generation results in the electrolyte solution of the potassium hydroxide series are shown in Fig. 8. 1 in the figure is a case in which the zinc powder having a concentration of iron of 5 ppm was used, and the hydrogen generation amount is generally large as compared with other two ones. 2 is a case in which the zinc powder having a concentration of iron of 4 ppm was used, and there is an effect of suppressing hydrogen generation at an adding amount of lead monoxide of not less than 10 ppm. 3 is a case in which the zinc powder having a concentration of iron of 2 ppm was used, and there is an effect in a concentration range equivalent to that of 2. Especially, when iron as an impurity was made not more than 4 ppm, the effect of the adding amount of lead monoxide of not more than 100 ppm is evident, and it is

possible to reduce the adding amount of lead as an environmental pollution substance.

The experiment has been described in the present Embodiment in which iron removal was performed by the magnet, and the concentration of iron is lowered, however, also in an experiment regarding zinc which was removed of iron by means of purification during production steps for zinc powder, substantially
5 the same results were obtained.

(Embodiment 16)

10 Into a positive electrode can were added a part of an electrolyte solution and 116 mg of a pellet (silver oxide content of 98 %) which was molded by adding a combined agent into silver oxide, and a separator of polyethylene and a separator of cellophane were set place. Next, a gasket of nylon was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, 30 mg of zinc powder, an inhibitor and the like were added, the remainder of the electrolyte solution was added dropwise, and then a negative
15 silver oxide batteries. The battery size made as SR621 type. The electrolyte solution was made again so that the adding amount of lead monoxide to be an amount with respect to the zinc amount.

They were stored at 60 °C for 20 days which is said to correspond to 1 year at ordinary temperature, and the self-discharge rate was measured, results of which are shown in Table 8.

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Table 8. Self-discharge rate of cell and hydrogen evolution rate

Experi- mental No.	Electro- lyte	Content of Fe in zinc powder ppm	Concentration of PbO in electrolyte ppm	Self- discharge rate %	Hydrogen evolution rate $\mu\text{L/g/day}$
1	KOH	5	0	13.2	0.98
2	KOH	5	10	8.7	0.76
3	KOH	5	50	7.2	0.79
4	KOH	5	100	4.0	0.48
5	KOH	5	500	8.6	0.64
6	KOH	5	1000	8.5	0.77
7	KOH	5	5000	16.2	1.48
8	KOH	4	0	13.0	0.89
9	KOH	4	10	3.8	0.45
10	KOH	4	50	3.5	0.37
11	KOH	4	100	2.2	0.33
12	KOH	4	500	2.8	0.29
13	KOH	4	1000	3.9	0.40
14	KOH	4	5000	17.6	1.42
15	KOH	2	0	11.0	0.82
16	KOH	2	10	2.2	0.24
17	KOH	2	50	1.5	0.23
18	KOH	2	100	1.4	0.21
19	KOH	2	500	2.2	0.25
20	KOH	2	1000	3.5	0.33
21	KOH	2	5000	17.2	1.62
22	NaOH	5	0	10.1	0.92
23	NaOH	5	10	8.0	0.72
24	NaOH	5	50	5.5	0.67
25	NaOH	5	100	4.1	0.40
26	NaOH	5	500	7.4	0.65
27	NaOH	5	1000	7.2	0.73
28	NaOH	5	5000	14.2	1.28
29	NaOH	4	0	9.2	0.80
30	NaOH	4	10	3.7	0.35
31	NaOH	4	50	2.0	0.29
32	NaOH	4	100	2.0	0.29
33	NaOH	4	500	2.2	0.26
34	NaOH	4	1000	3.2	0.41
35	NaOH	4	5000	15.2	1.31
36	NaOH	2	0	7.1	0.72
37	NaOH	2	10	1.5	0.25
38	NaOH	2	50	0.8	0.19
39	NaOH	2	100	0.7	0.18
40	NaOH	2	500	1.1	0.23
41	NaOH	2	1000	2.9	0.28
42	NaOH	2	5000	9.8	1.46

As seen also from the results, it is understood that the self-discharge rate can be suppressed to be not more than 4 % by adding 10-1000 ppm of lead monoxide in the battery using the zinc powder having iron of not more than 4 ppm. The decrease in the battery capacity by the addition of lead monoxide at the high concentration is considered if it is due to the fact that needle-like crystals of lead grew on account of excessive addition, and a short circuit might be caused.

(Embodiment 17)

A button type silver oxide battery was manufactured in the same manner as Embodiment 16 except that a specially prepared gasket of polypropylene for allowing hydrogen to penetrate was used. Ten pieces of

the manufactured silver oxide batteries were put into vessel made of glass filled with liquid paraffin in a high temperature tank, a collecting tube having graduations at the upper portion was attached, and the amount of generated hydrogen gas was measured. This state was maintained at 60 °C for 20 days which is said to correspond to a term of about 1 year, and the hydrogen gas generation amount after 20 days was investigated. Results are shown in Table 8.

As understood from Embodiments, the hydrogen generation is mild and the self-discharge is also less in the case of the sodium hydroxide series as compared with the potassium hydroxide series with respect to the electrolyte solution.

In addition, the less the iron content in the zinc powder is, the better results are obtained, and the smaller amount the adding amount of lead monoxide is.

In the actual trial production of batteries, there is given a value smaller than the hydrogen gas generation amount in the experiment in Embodiment 15. In this fact, it is considered that the copper piece used in Embodiment 15 is different from a structure of an actual collector, a part of generated hydrogen gas has been consumed by the reduction of silver oxide and the like.

The method in Embodiment 15 is different in the amount of order of hydrogen gas generation, but it is sufficient as a method for observing substituted characteristics for predicting hydrogen generation in an actual battery.

The same experiment was performed for indium compounds, tin compounds containing tetravalent tin, and mixtures thereof other than lead monoxide. Results thereof are substantially the same as those in the case of lead monoxide, and it has been found that the self-discharge rate can be suppressed to be not more than 4 % by adding 10-1000 ppm of the inhibitor.

In the present invention, the description has been made for iron which is easiest to contaminate and has a high risk of hydrogen generation, however, it is needless to say that it is preferable one in which impurities such as nickel, cobalt, antimony and the like reduced as far as possible. In addition, in the case of the negative electrode active material such as zinc and the like in which hydrogen generation is large, a homogeneous film is difficult to be formed due to the hydrogen generation which occurs as a competitive reaction against lead deposition onto the surface, and the effect of lead deposition becomes small. Thus, it is more effective to use a negative electrode active material in which zinc is added with indium, bismuth, lead, aluminum, gallium, calcium and the like so as to suppress the hydrogen generation to some extent.

In accordance with Embodiments 18-21, the effect of the case in which the moisture in the battery is controlled will be explained.

(Embodiment 18)

Into a positive electrode can were added a part of an electrolyte solution and a pellet (silver oxide content of 98 %) which was molded by putting a combined agent into silver oxide, and a separator of polyethylene and a separator of cellophane were set place.

Next, a gasket of specially prepared polypropylene for allowing hydrogen to penetrate was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, zinc powder, an inhibitor and the like were added, the remainder of the electrolyte solution was added dropwise, and then a negative electrode can was set place to perform sealing to manufacture a button type silver oxide battery.

The electrolyte solution was prepared such that by making a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide series, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide series, as a base, and the inhibitor was optionally added thereto. When the inhibitor is added, its amount was made as 1000 ppm with respect to the zinc weight, and it was dissolved in the electrolyte solution.

Ten pieces of the manufactured coin or button silver oxide batteries were put into a vessel made of glass filled with liquid paraffin in a high temperature tank, a collecting tube having graduations at the upper portion was attached, and the amount of generated hydrogen was measured. In this state, it was maintained at 60 °C for 20 days, which is said to correspond to a term of about 1 year, and the hydrogen gas generation amount and the self-discharge rate after 20 days were investigated. The battery capacity was measured by connecting a resistance of 25 Ω and having it discharge. The self-discharge rate was calculated from the change in the battery capacity before and after the storage at 60 °C for 20 days regarding the same kind of batteries.

The size of the coin or button silver oxide battery manufactured and evaluated, the content of metal in the zinc powder used, the electrolyte solution, the moisture amount, the inhibitor added, and the hydrogen generation amount and the self-discharge rate after the storage at 60 °C for 20 days are shown in Table 9

and Table 10.

Table 9. Hydrogen evolution rate and self-discharge rate of cell

Experi- mental No.	Cell type	Composition of added metal (ppm vs zinc)					Electro- lyte	Amount of water added to zinc of 1 mg	Inhibitor	After 20 days storage at 60 °C		After 1 year storage at room temperature	
		Ga	In	Pb	Bi	Al				Hydrogen evolution rate $\mu\text{l/g/day}$	Self- discharge rate %	Hydrogen evolution rate $\mu\text{l/g/day}$	Self- discharge rate %
1	SR628		500	500	500		NaOH	0.448	—	0.37	3.00	0.00	3.10
2	SR628		500	500	500		KOH	0.428	—	0.39	2.50	0.00	2.50
3	SR628			500	500		NaOH	0.448	—	0.41	2.70	0.00	2.80
4	SR628			500	500		KOH	0.428	—	0.47	2.40	0.00	2.50
5	SR628	100	200	500		450	NaOH	0.448	—	0.40	2.50	0.00	2.80
6	SR628	100	200	500		450	KOH	0.428	—	0.46	2.60	0.00	2.80
7	SR621		500	500	500		NaOH	0.459	—	0.38	2.10	0.00	2.10
8	SR621		500	500	500		KOH	0.440	—	0.40	3.10	0.00	3.30
9	SR621			500	500		NaOH	0.459	—	0.43	2.80	0.00	2.70
10	SR621			500	500		KOH	0.440	—	0.45	2.70	0.00	2.70
11	SR516		500	500	500		NaOH	0.476	—	0.35	2.80	0.00	2.70
12	SR516		500	500	500		KOH	0.455	—	0.37	2.80	0.00	2.80
13	SR516			500	500		NaOH	0.476	—	0.41	2.90	0.00	2.50
14	SR516			500	500		KOH	0.455	—	0.45	3.00	0.00	2.90
15	SR527		500	500	500		NaOH	0.488	—	0.38	2.70	0.00	2.80
16	SR527		500	500	500		KOH	0.463	—	0.39	2.80	0.00	2.60
17	SR527			500	500		NaOH	0.488	—	0.43	2.50	0.00	2.40
18	SR527			500	500		KOH	0.463	—	0.47	2.70	0.00	2.60
19	SR726		500	500	500		NaOH	0.441	—	0.35	2.10	0.00	2.50

Table 10. Hydrogen evolution rate and self-discharge rate of cell

Experi- mental No.	Cell type	Composition of added metal (ppm vs zinc)					Electro- lyte	Amount of water added to zinc of 1 mg	Inhibitor	After 20 days storage at 60 °C		After 1 year storage at room temperature	
		Ga	In	Pb	Bi	Al				Hydrogen evolution rate $\mu\text{l/g/day}$	Self- discharge rate %	Hydrogen evolution rate $\mu\text{l/g/day}$	Self- discharge rate %
20	SR728		500	500	500		KOH	0.423	—	0.38	1.50	0.00	2.00
21	SR726			500	500		NaOH	0.441	—	0.37	1.80	0.00	1.80
22	SR728			500	500		KOH	0.423	—	0.45	2.20	0.00	2.10
23	SR1120		500	500	500		NaOH	0.441	—	0.35	1.90	0.00	2.10
24	SR1120		500	500	500		KOH	0.423	—	0.34	2.80	0.00	2.10
25	SR1120			500	500		NaOH	0.441	—	0.37	2.30	0.00	2.20
26	SR1120			500	500		KOH	0.423	—	0.38	2.40	0.00	2.50
27	SR028		500	500	500		NaOH	0.448	Indium sulfate	0.33	2.50	0.00	2.50
28	SR028		500	500	500		KOH	0.428	Indium sulfate	0.35	2.30	0.00	2.60
29	SR028			500	500		NaOH	0.448	Indium sulfate	0.38	2.70	0.00	2.50
30	SR028			500	500		KOH	0.428	Indium sulfate	0.42	2.80	0.00	2.90
31	SR028	100	200	500		450	NaOH	0.448	Indium sulfate	0.37	2.70	0.00	2.70
32	SR028	100	200	500		450	KOH	0.428	Indium sulfate	0.41	2.80	0.00	2.80
33	SR028			500			NaOH	0.448	Indium sulfate	0.39	2.80	0.00	2.70
34	SR028			500			KOH	0.428	Indium sulfate	0.42	2.60	0.00	2.80
35	SR028			500			NaOH	0.448	Indium sulfamate	0.42	2.70	0.00	2.50
36	SR028			500			KOH	0.428	Indium sulfamate	0.45	2.70	0.00	2.40
37	SR028			500			NaOH	0.448	PbO	0.42	2.50	0.00	2.90
38	SR028			500			KOH	0.428	PbO	0.44	2.50	0.00	2.20

In Table 9 and Table 10, with respect to the added metal composition, the weight for zinc was expressed by ppm. The moisture amount was represented by how much mg of water was contained with respect to 1 mg of zinc. The unit for the hydrogen generation amount is $\mu\text{l/g/day}$, the hydrogen generation amount per one day with respect to the zinc amount is shown at the right of each of the columns for 60 °C for 20 days and ordinary temperature for 1 year in the table, and the unit for the self-discharge rate is

shown by %. In-sulfate indicates indium sulfate, In-sulfamate indicates indium sulfamate, and PbO indicates lead oxide. It is understood that all are suppressed to be not more than $0.03 \mu \text{L/g/day}$ when the hydrogen generation amount is calculated into a value at ordinary temperature. In addition, it was also possible to make the self-discharge rate to be not more than 4 %/year.

5
(Embodiment 19)

In order to know whether or not the storage at 60°C for 20 days in Embodiment 18 corresponds to 1 year at ordinary temperature, a test for 1 year was actually performed.

10 The gasket used in Embodiment 18 was exchanged into one made of nylon being used in ordinary production of silver oxide batteries, and coin or button silver oxide batteries were manufactured in the same manner. Ten pieces of the manufactured silver oxide batteries were put into a vessel made of glass filled with liquid paraffin in a high temperature tank maintained at 25°C , and a collecting tube for generating hydrogen gas was attached at the upper portion. This state was maintained for 1 year, and the hydrogen
15 generation amount and the self-discharge rate after 1 year were investigated. The measurement method was the same as that in Embodiment 18. Results of the hydrogen generation amount and the self-discharge rate are shown at the right ends in Table 9 and Table 10.

As a result of the test, no gas generation was observed in any sample. This is considered to be due to the fact that a minute amount of hydrogen generated in the battery was consumed by the reduction of silver
20 oxide. The self-discharge rate become substantially the same as those preserved at 60°C for 20 days, and all had values of not more than 4 %.

(Embodiment 20)

25 As for the electrolyte solution, in the case of the potassium hydroxide series, one in which zinc oxide was added to potassium hydroxide up to approximate saturation was used. The amount of the electrolyte solution to be added per 1 mg of zinc was changed from 0.37 to 0.96 mg which was from 0.25 to 0.65 mg as calculated into the amount of water, using zinc containing 500 ppm of each of indium, lead and bismuth and a gasket of polypropylene in the same manner as Embodiment 18, a silver oxide battery of the SR626
30 size was made, and the storage test was performed at 60°C for 20 days.

The hydrogen generation amount with respect to the added amount of water is shown in Fig. 9. The hydrogen generation amount indicates an amount generated from 1 g of zinc per one day, and the unit is $\mu \text{L/g/day}$. According to the result, it is understood that when the amount of water is not less than 0.31 mg, the hydrogen generation amount is lowered to not more than $0.54 \mu \text{L/g/day}$ ($0.03 \mu \text{L/g/day}$ as calculated
35 into a value at ordinary temperature). In addition, when the amount of water was not less than 0.57 mg, the occurrence of liquid leakage was observed.

In the same manner, an electrolyte solution of the sodium hydroxide series was used, the amount of the electrolyte solution to be added per 1 mg of zinc was changed from 0.36 to 0.92 mg which is from 0.25 to 0.65 mg when it is calculated into the amount of water, and the storage test was performed at 60°C for 20
40 days. A result is shown in Fig. 10. It is understood that when the amount of water is not less than 0.32 mg, the hydrogen generation amount is lowered to not more than $0.54 \mu \text{L/g/day}$. In addition, when the amount of water became not less than 0.59 mg, the occurrence of liquid leakage was observed.

Namely, it is desirable that the amount of water to be added per 1 mg of zinc is 0.31-0.57 mg when the electrolyte solution is the potassium hydroxide series, and it is 0.32-0.59 mg in the case of sodium
45 hydroxide.

(Embodiment 21)

In order to know whether or not there is actually a problem in the range of the amount of water in which
50 the hydrogen generation is not more than $0.54 \mu \text{L/g/day}$ and there is no liquid leakage in Embodiment 20, the storage test at ordinary temperature for 1 year was performed and tried using the method in Embodiment 19. In the case of the potassium hydroxide series, the electrolyte solution was added as that the amount of water to be added per 1 mg of zinc became 0.31, 0.44 and 0.57 mg. In the same manner, in the case of the sodium hydroxide series, the electrolyte solution was added so that the amount of water
55 became 0.32, 0.45 and 0.59 mg. The gasket was changed into nylon in the same manner as Embodiment 20, and 10 pieces of silver oxide batteries were manufactured, respectively. According to the results of the storage test at ordinary temperature for 1 year for the manufactured silver oxide batteries, there was no hydrogen generation, the self-discharge rates were also not more than 4 % for all ones, and it has been

found that there is no practical problem.

In accordance with Embodiments 22-24, the effect of the case in which each technique is combined will be explained.

Before actually investigating characteristics with batteries, a specially prepared test tube was used to investigate how much amount of hydrogen gas was generated from each of inhibitors and plated copper plates. The method therefor will be described hereinafter.

Into the specially prepared test tube having a volume of 25 ml and graduated to know the gas generation amount were put beforehand 2 g of zinc powder containing 500 ppm of each of bismuth, indium and lead manufactured by the atomization method, and a copper piece as the same material as a collector having an area of 0.6 cm² and a thickness of 0.1 mm, an electrolyte solution to be tested was added thereto to heat to 60 °C, and the volume of generated hydrogen gas was measured for 7 days. The number of repetition of the test was 10, and an average value thereof was used as a result. The electrolyte solution was prepared such that by making a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide series, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide series as a base, and an inhibitor was optionally added thereto.

(Embodiment 22)

The hydrogen generation from zinc was measured by the combination of a copper plate coated by the electric plating and the substitution plating and the inhibitor. The plating treatment is shown hereinafter.

* Indium (In) plating

method: electric plating
 plating bath: 25 °C
 : indium sulfate 60 g/l
 : sodium sulfate 10 g/l
 plating film thickness: 0.3 μm

* Tin (Sn) plating

method: electric plating
 plating bath: 70 °C
 : potassium stannate 100 g/l
 : potassium hydroxide 15 g/l
 : potassium acetate 5 g/l
 plating film thickness: 0.3 μm

* Zinc indium (Zn-In) plating

method: contact plating (copper plate and zinc are put into the following plating bath heated to 60 °C and left for 1 hour)
 plating bath: KOH 30 %
 : ZnO saturation
 : indium sulfate 0.3 %
 plating film thickness: 0.3 μm

The plating onto the copper plate and the hydrogen gas generation measurement result with respect to the inhibitor are shown as the case using the electrolyte solution of the KOH series in Table 11 and that of the NaOH type in Table 12.

Table 11

Hydrogen evolution rate with plated copper plate				
Experimental No.	Kind of plating	Electrolyte	Inhibitor (concentration: 100ppm)	Hydrogen evolution rate μ L/g/day
1	In	KOH	-	31.43
2	In	KOH	Indium sulfate	14.28
3	In	KOH	Indium sulfamate	12.41
4	In	KOH	Indium hydroxide	16.73
5	In	KOH	Lead oxide	17.85
6	In	KOH	Barium hydroxide	8.57
7	In	KOH	Carbon fluoride • POE	12.14
8	Sn	KOH	-	35.78
9	Sn	KOH	Indium sulfate	16.11
10	Sn	KOH	Indium sulfamate	17.72
11	Sn	KOH	Indium hydroxide	16.53
12	Sn	KOH	Lead oxide	17.91
13	Sn	KOH	Barium hydroxide	10.89
14	Sn	KOH	Carbon fluoride • POE	14.38
15	Zn • In	KOH	-	32.34
16	Zn • In	KOH	Indium sulfate	15.27
17	Zn • In	KOH	Indium sulfamate	13.35
18	Zn • In	KOH	Indium hydroxide	18.83
19	Zn • In	KOH	Lead oxide	16.82
20	Zn • In	KOH	Barium hydroxide	11.30
21	Zn • In	KOH	Carbon fluoride • POE	13.01

Table 12

Hydrogen evolution rate with plated copper plate				
Experimental No.	Kind of plating	Electrolyte	Inhibitor (concentration: 100ppm)	Hydrogen evolution rate μ L/g/day
22	In	NaOH	-	29.41
23	In	NaOH	Indium sulfate	14.75
24	In	NaOH	Indium sulfamate	10.58
25	In	NaOH	Indium hydroxide	15.37
26	In	NaOH	Lead oxide	14.14
27	In	NaOH	Barium hydroxide	7.82
28	In	NaOH	Carbon fluoride-POE	9.34
29	Sn	NaOH	-	30.10
30	Sn	NaOH	Indium sulfate	15.03
31	Sn	NaOH	Indium sulfamate	15.28
32	Sn	NaOH	Indium hydroxide	15.38
33	Sn	NaOH	Lead oxide	14.53
34	Sn	NaOH	Barium hydroxide	8.72
35	Sn	NaOH	Carbon fluoride-POE	12.43
36	Zn-In	NaOH	-	32.83
37	Zn-In	NaOH	Indium sulfate	14.98
38	Zn-In	NaOH	Indium sulfamate	13.19
39	Zn-In	NaOH	Indium hydroxide	14.13
40	Zn-In	NaOH	Lead oxide	14.27
41	Zn-In	NaOH	Barium hydroxide	9.98
42	Zn-In	NaOH	Carbon fluoride-POE	9.84

The concentration of the inhibitor is shown by a value with respect to the electrolyte solution. As the surfactant of the fluorocarbon(carbon fluoride)-polyoxyethylene (POE) series, one having a carbon number of fluorocarbon of 10 and a polymerization degree of POE of 50 was used. Usually, the same effect can be obtained if the one having a carbon number of fluorocarbon of 4-20 and a polymerization degree of POE of 30-100 is used.

As compared with the one in which no inhibitor is added, in the one in which the inhibitor of the present invention is added, the hydrogen gas generation amount decreases to about 1/3 to 1/2, and it is understood that there is an effect in relation to suppression of the hydrogen gas generation. The difference due to the plating or the inhibitor is not presented so much. In addition, the hydrogen gas generation amount with respect to the number of test days is shown in Fig. 11 as for the measurement result of the case using indium plated copper plates. As for the indium compound, only indium sulfate is shown in the figure.

In the present Embodiment, the result of the case using one having a plating film thickness of $0.3 \mu m$ has been shown, however, in practice, also when one having a film thickness of $0.1-1 \mu m$ was used, there was little difference in the result.

(Comparative example 2)

For comparison, an experiment was performed using non-plated copper plates in the same procedure as Embodiment 22. Experiment results are shown in Table 13 and Fig. 12.

Table 13

Hydrogen evolution rate with not plated copper plate			
Example No.	Electrolyte	Inhibitor (concentration: 100ppm)	Hydrogen evolution rate μ L/g/day
1	KOH	-	516.00
2	KOH	Indium sulfate	30.56
3	KOH	Indium sulfamate	28.73
4	KOH	Indium hydroxide	29.07
5	KOH	Lead oxide	24.15
6	KOH	Barium hydroxide	480.32
7	KOH	Carbon fluoride • POE	465.29
8	NaOH	-	450.24
9	NaOH	Indium sulfate	25.86
10	NaOH	Indium sulfamate	27.59
11	NaOH	Indium hydroxide	25.48
12	NaOH	Lead oxide	20.25
13	NaOH	Barium hydroxide	400.44
14	NaOH	Carbon fluoride • POE	392.91

It is understood that in the case of no addition of the inhibitor and addition of barium hydroxide or the surfactant of the fluorocarbon-polyoxyethylene series, the hydrogen gas generation amount increases with respect to the number of test days exponentially. On the other hand, it is understood that in the case of addition of indium sulfate and lead oxide, the generation of hydrogen gas is suppressed. This is due to the fact that the compound of indium or lead has the function to coat the copper plate, but barium hydroxide and the surfactant have no such a function.

(Embodiment 23)

Using a copper plate plated in the same manner as Embodiment 22, the hydrogen gas generation in the case where it is used by combining inhibitors was measured using an electrolyte solution of the KOH series. Results are shown in Table 14.

Table 14

Hydrogen evolution rate with indium plated copper plate and inhibitor					
Experimental No.	Concentration of inhibitor ppm				Hydrogen evolution rate μ L/g/day
	Indium sulfate	Lead oxide	Indium hydroxide	Carbon fluoride-POE	
43	100	100	0	0	15.82
44	100	0	100	0	10.10
45	100	0	0	100	13.41
46	0	100	100	0	14.64
47	0	100	0	100	15.54
48	0	0	100	100	12.47
49	100	100	100	0	14.87
50	100	100	0	100	12.21
51	100	0	100	100	13.05
52	100	100	100	100	13.94

In the column for the inhibitor in the table, the concentration of each inhibitor added to the electrolyte solution with respect to the electrolyte solution is shown.

It is understood that the hydrogen gas generation is in a low level similar as the Embodiments, and there is not too much bad effect even when a plurality of inhibitors are used.

Next, a battery was actually manufactured, and the effects of the plating on the collector and the inhibitor were investigated.

(Embodiment 24)

Into a positive electrode can were added a part of an electrolyte solution and 116 mg of a pellet (silver oxide content of 98 %) was molded by putting a combined agent into which silver oxide, and a separator of polyethylene and a separator of cellophane were set place. Next, a gasket of nylon was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, and 30 mg of zinc powder were added, the remainder of the electrolyte solution optionally added with an inhibitor was added dropwise, and then a negative electrode can was set place to perform sealing to manufacture 100 pieces for every kind of button type silver oxide batteries.

The type and the film thickness of plating onto the negative electrode can, the type and the concentration of the added inhibitor, the closed circuit voltage and the self-discharge rate are shown in Tables 15 and 16.

Table 15. Characteristics of cell with KOH electrolyte.

Experi- mental No.	Kind of plating	Concentration of inhibitor ppm				Self-discharge rate %		Closed circuit voltage V	
		Indium sulfate	Lead oxide	barium hydroxide	Carbon fluoride POE	Fresh cell	A part- discharged cell	Fresh cell	A part- discharged cell
53	-	0	0	0	0	5.6	9.4	1.159	1.180
54	In	0	0	0	0	4.5	6.2	1.165	1.192
55	In	1000	0	0	0	2.6	2.7	1.169	1.198
56	In	0	1000	0	0	2.7	2.8	1.168	1.197
57	In	0	0	1000	0	2.5	4.9	1.185	1.198
58	In	0	0	0	1000	2.9	5.1	1.165	1.185
59	In	1000	1000	0	0	2.5	2.6	1.168	1.198
60	In	1000	0	1000	0	2.4	2.6	1.185	1.239
61	In	1000	0	0	1000	2.5	2.7	1.168	1.198
62	In	0	1000	0	1000	2.6	2.7	1.167	1.197
63	In	0	1000	1000	0	2.7	2.7	1.167	1.238
64	In	0	1000	1000	1000	2.6	2.6	1.186	1.238
65	In	1000	1000	1000	0	2.6	2.7	1.186	1.129
66	In	1000	1000	0	1000	2.7	2.7	1.168	1.198
67	In	1000	0	1000	1000	2.6	2.6	1.185	1.238
68	In	1000	1000	1000	1000	2.6	2.7	1.185	1.238
69	Sn	0	0	0	0	5.7	10.5	1.160	1.181
70	Sn	1000	0	0	0	4.7	6.8	1.166	1.193
71	Sn	0	1000	0	0	2.7	2.7	1.171	1.198
72	Sn	0	0	1000	0	2.7	2.8	1.170	1.198
73	Sn	0	0	0	1000	2.6	5.0	1.186	1.197
74	Sn	1000	1000	0	0	3.0	5.3	1.165	1.187
75	Sn	1000	0	1000	0	2.6	2.7	1.169	1.199
76	Sn	1000	0	0	1000	2.4	2.5	1.187	1.240
77	Sn	0	1000	0	1000	2.4	2.8	1.169	1.200
78	Sn	0	1000	1000	0	2.5	2.6	1.168	1.197
79	Sn	0	1000	1000	1000	2.7	2.8	1.169	1.240
80	Sn	1000	1000	1000	0	2.5	2.8	1.188	1.240
81	Sn	1000	1000	0	1000	2.8	2.8	1.186	1.131
82	Sn	1000	0	1000	1000	2.8	2.8	1.170	1.199
83	Sn	1000	1000	1000	1000	2.7	2.6	1.186	1.239
84	Zn-In	0	0	0	0	5.7	9.8	1.166	1.193
85	Zn-In	1000	0	0	0	4.9	7.1	1.170	1.199
86	Zn-In	0	1000	0	0	2.8	2.7	1.170	1.197
87	Zn-In	0	0	1000	0	2.9	2.9	1.187	1.199
88	Zn-In	0	0	0	1000	2.6	4.8	1.166	1.184
89	Zn-In	1000	1000	0	0	2.9	5.3	1.168	1.200
90	Zn-In	1000	0	1000	0	2.6	2.7	1.186	1.240
91	Zn-In	1000	0	0	1000	2.6	2.7	1.170	1.199
92	Zn-In	0	1000	0	1000	2.6	2.9	1.168	1.199
93	Zn-In	0	1000	1000	0	2.7	2.7	1.168	1.238
94	Zn-In	0	1000	1000	1000	2.9	2.9	1.188	1.240
95	Zn-In	1000	1000	1000	0	2.8	2.8	1.188	1.131
96	Zn-In	1000	1000	0	1000	2.6	2.9	1.168	1.200
97	Zn-In	1000	0	1000	1000	2.9	2.8	1.187	1.239
98	Zn-In	1000	1000	1000	1000	2.7	2.7	1.186	1.239

Table 16. Characteristics of cell with NaOH electrolyte

Experimental No.	Kind of plating	Concentration of inhibitor ppm				Self-discharge rate %		Closed circuit voltage V	
		Indium sulfate	Lead oxide	barium hydroxide	Carbon fluoride · POE	Fresh cell	A part-discharged cell	Fresh cell	A part-discharged cell
99	—	0	0	0	0	5.3	9.0	1.158	1.180
100	In	0	0	0	0	4.2	5.9	1.164	1.193
101	In	1000	0	0	0	2.2	2.3	1.169	1.199
102	In	0	1000	0	0	2.2	2.4	1.167	1.197
103	In	0	0	1000	0	2.2	4.6	1.187	1.199
104	In	0	0	0	1000	2.5	4.8	1.166	1.186
105	In	1000	1000	0	0	2.1	2.3	1.169	1.199
106	In	1000	0	1000	0	1.9	2.2	1.186	1.239
107	In	1000	0	0	1000	2.0	2.5	1.169	1.197
108	In	0	1000	0	1000	2.3	2.5	1.169	1.196
109	In	0	1000	1000	0	2.3	2.2	1.169	1.238
110	In	0	1000	1000	1000	2.0	2.1	1.187	1.237
111	In	1000	1000	1000	0	2.4	2.4	1.186	1.131
112	In	1000	1000	0	1000	2.4	2.3	1.170	1.198
113	In	1000	0	1000	1000	2.3	2.0	1.187	1.239
114	In	1000	1000	1000	1000	2.2	2.5	1.186	1.237
115	Sn	0	0	0	0	5.5	10.2	1.160	1.183
116	Sn	1000	0	0	0	4.4	6.5	1.167	1.194
117	Sn	0	1000	0	0	2.4	2.3	1.173	1.199
118	Sn	0	0	1000	0	2.5	2.6	1.171	1.200
119	Sn	0	0	0	1000	2.4	4.7	1.186	1.197
120	Sn	1000	1000	0	0	2.7	5.0	1.166	1.186
121	Sn	1000	0	1000	0	2.2	2.5	1.170	1.201
122	Sn	1000	0	0	1000	2.1	2.3	1.187	1.241
123	Sn	0	1000	0	1000	2.2	2.5	1.170	1.201
124	Sn	0	1000	1000	0	2.2	2.2	1.167	1.198
125	Sn	0	1000	1000	1000	2.2	2.5	1.171	1.241
126	Sn	1000	1000	1000	0	2.3	2.6	1.189	1.242
127	Sn	1000	1000	0	1000	2.6	2.5	1.187	1.133
128	Sn	1000	0	1000	1000	2.6	2.3	1.172	1.200
129	Sn	1000	1000	1000	1000	2.4	2.4	1.186	1.239
130	Zn·In	0	0	0	0	5.3	9.5	1.168	1.194
131	Zn·In	1000	0	0	0	4.6	6.6	1.171	1.201
132	Zn·In	0	1000	0	0	2.4	2.5	1.172	1.198
133	Zn·In	0	0	1000	0	2.5	2.7	1.187	1.200
134	Zn·In	0	0	0	1000	2.3	4.6	1.168	1.185
135	Zn·In	1000	1000	0	0	2.6	5.0	1.170	1.200
136	Zn·In	1000	0	1000	0	2.3	2.3	1.188	1.241
137	Zn·In	1000	0	0	1000	2.2	2.4	1.171	1.198
138	Zn·In	0	1000	0	1000	2.4	2.5	1.169	1.199
139	Zn·In	0	1000	1000	0	2.6	2.5	1.167	1.237
140	Zn·In	0	1000	1000	1000	2.7	2.6	1.187	1.242
141	Zn·In	1000	1000	1000	0	2.4	2.3	1.188	1.132
142	Zn·In	1000	1000	0	1000	2.6	2.7	1.167	1.201
143	Zn·In	1000	0	1000	1000	2.6	2.6	1.189	1.240
144	Zn·In	1000	1000	1000	1000	2.5	2.5	1.186	1.240

The inhibitor concentration in the table indicates a value with respect to the zinc weight. The measurement of the self-discharge rate was performed after maintaining it at 60 °C for 20 days, which is said to correspond to 1 year.

The self-discharge rate after partial discharge was measured by being left at 60 °C for 20 days after 50 % depth discharge (partial discharge). The closed circuit voltage was measured at -10 °C before discharge and after partial discharge.

As compared with those of no addition of inhibitors (Embodiments No. 54, 69 and 84 in Table 15 and Embodiments No. 100, 115 and 130 in Table 16), those added with inhibitors have small self-discharge rates. With respect to the partial discharge, in the case of those added with one of the type for coating the negative electrode can which being copper such as indium sulfate and lead monoxide, the self-discharge rate is improved. It is understood that the closed circuit voltage becomes high in those added with barium hydroxide. It is understood from the table that when the inhibitor of the coating type and barium hydroxide are added, the self-discharge rate before discharge, the self-discharge rate after partial discharge, and the closed circuit voltage are improved. In the present experiment, a behavior that the closed circuit voltage after partial discharge becomes higher than that before discharge has been presented which is opposite to a case using mercurated zinc. The cause thereof is now under investigation.

The plating onto the negative electrode can was performed in the same manner as Embodiment 22, and results for each film thickness of $0.3 \mu\text{m}$ have been shown. There was little difference in the result even when those having a film thickness of $0.1\text{--}1 \mu\text{m}$ were used.

Further, with a battery manufactured using an electrolyte of the KOH type and a negative electrode can applied with indium plating, the self-discharge rate after lapse of corresponding 1 year and the self-discharge rate after partial discharge with respect to the inhibitor concentration were measured.

Results are shown in Figs. 13 and 14. Provided that the practical range is not more than about 3 % of the self-discharge rate, it has been found that the effect is provided for indium sulfate of 50-5000 ppm, for lead monoxide of 20-5000 ppm, for barium hydroxide of not less than 50 ppm, and for the fluorocarbon-polyoxyethylene type surfactant of not less than 100 ppm as the concentration with respect to zinc. With respect to the self-discharge rate after 50 % depth discharge, the effect was provided for indium sulfate of 50-5000 ppm, and for lead monoxide of 20-5000 ppm. The effect is not provided so much after partial discharge for barium hydroxide, and the fluorocarbon-polyoxyethylene series surfactant. The result has been shown in the case of indium sulfate for indium compounds, however, the effect was also provided for other compounds in substantially the same concentration range. In addition, even when an electrolyte solution of the NaOH series and other negative electrode can plating were used, the concentration range of the inhibitor exhibiting the effect was substantially the same.

In accordance with Embodiments 25-29, the effect of the case combined each techniques will be explained. In Embodiments 22-24, the evaluation was made using combinations in which the surfactants were changed. In the trial production of batteries, the evaluation was made also by changing the type of zinc.

Before actually investigating characteristics using batteries, a specially prepared test tube was used to investigate how much amount of hydrogen gas was generated from each of inhibitors and plated copper plates. The method therefor will be described hereinafter.

Into the specially prepared test tube having a volume of 25 ml and graduated to know the gas generation amount were added beforehand 2 g of zinc powder containing 500 ppm of each of bismuth, indium and lead manufactured by the atomization method, and a copper piece as the same material as a collector having an area of 0.6 cm^2 and a thickness of 0.1 mm , an electrolyte solution to be tested was added thereto to heat to 60°C , and the volume of generated hydrogen gas was measured for 7 days. The number of repetition of the test was 10, and an average value thereof was used as a result. The electrolyte solution was prepared such that by making a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide type, or a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide type, as a haze and an inhibitor was optionally added thereto.

(Embodiment 25)

The hydrogen generation from zinc was measured by the combination of a copper plate coated by the electric plating and the substitution plating and the inhibitor. The plating treatment is shown hereinafter.

* Indium (In) plating

method: electric plating

plating bath: indium sulfate 60 g/l, sodium sulfate 10 g/l, 25°C

plating film thickness: $0.3 \mu\text{m}$

* Tin (Sn) plating

method: electric plating

plating bath: potassium stannate 100 g/l, potassium hydroxide 15 g/l, potassium acetate 5 g/l, 70 °C

5 plating film thickness: 0.3 μ m

* zinc indium (Zn-In) plating

10 method: contact plating; copper plate and zinc are added into the following plating bath heated to 60 °C and left for 1 hour

plating bath: KOH 30 %, ZnO saturation, indium sulfate 0.3 %

plating film thickness: 0.3 μ m

15 The plating onto the copper plate and the hydrogen gas generation measurement result with respect to the inhibitor are shown as the case using the electrolyte solution of the KOH series in Table 17, while that of the NaOH series in Table 18. The concentration of the inhibitor is shown by a value with respect to the electrolyte solution.

Table 17

Hydrogen evolution rate with plated copper plate				
Experimental No.	Kind of plating	Electrolyte	Inhibitor (concentration: 100ppm)	Hydrogen evolution rate μ L/g/day
1	In	KOH	-	31.43
2	In	KOH	Indium sulfate	14.28
3	In	KOH	Indium sulfamate	12.41
4	In	KOH	Indium hydroxide	16.73
5	In	KOH	Lead oxide	17.85
6	In	KOH	Barium hydroxide	8.57
7	In	KOH	POERA	8.42
8	Sn	KOH	-	35.78
9	Sn	KOH	Indium sulfate	16.11
10	Sn	KOH	Indium sulfamate	17.72
11	Sn	KOH	Indium hydroxide	16.53
12	Sn	KOH	Lead oxide	17.91
13	Sn	KOH	Barium hydroxide	10.89
14	Sn	KOH	POERA	10.02
15	Zn-In	KOH	-	32.34
16	Zn-In	KOH	Indium sulfate	15.27
17	Zn-In	KOH	Indium sulfamate	13.35
18	Zn-In	KOH	Indium hydroxide	18.83
19	Zn-In	KOH	Lead oxide	16.82
20	Zn-In	KOH	Barium hydroxide	11.30
21	Zn-In	KOH	POERA	10.98

Table 18

Hydrogen evolution rate with plated copper plate				
Experimental No.	Kind of plating	Electrolyte	Inhibitor (concentration: 100ppm)	Hydrogen evolution rate μ L/g/day
22	In	NaOH	-	29.41
23	In	NaOH	Indium sulfate	14.75
24	In	NaOH	Indium sulfamate	10.58
25	In	NaOH	Indium hydroxide	15.37
26	In	NaOH	Lead oxide	14.14
27	In	NaOH	Barium hydroxide	7.82
28	In	NaOH	POERA	8.13
29	Sn	NaOH	-	30.10
30	Sn	NaOH	Indium sulfate	15.03
31	Sn	NaOH	Indium sulfamate	15.28
32	Sn	NaOH	Indium hydroxide	15.38
33	Sn	NaOH	Lead oxide	14.53
34	Sn	NaOH	Barium hydroxide	8.72
35	Sn	NaOH	POERA	9.52
36	Zn-In	NaOH	-	32.83
37	Zn-In	NaOH	Indium sulfate	14.98
38	Zn-In	NaOH	Indium sulfamate	13.19
39	Zn-In	NaOH	Indium hydroxide	14.13
40	Zn-In	NaOH	Lead oxide	14.27
41	Zn-In	NaOH	Barium hydroxide	9.98
42	Zn-In	NaOH	POERA	8.95

Polyoxyethylene alkylamide is the one (represented by POERA in the table and the figure), in which two polyoxyethylenes (POE) are bonded to the alkyl group at nitrogen through amide bond. In this case, the one having a carbon number of the alkyl group of 11 and a polymerization degree of POE of 15 was used. Usually, when the one having a carbon number of the alkyl group of 3-30 and a polymerization degree of POE of 2-50 is used, the same effect is obtained.

As compared with the one in which no inhibitor is added, in the case of the one in which the inhibitor of the present invention is added, the hydrogen gas generation amount decreases to about 1/3 to 1/2, and it is understood that there is an effect in relation to suppression of the hydrogen gas generation. The difference due to the plating or the inhibitor is not presented so much. In addition, the hydrogen gas generation amount with respect to the number of test days is shown in Fig. 15 as the measurement result of the case using indium plated copper plates. As for the indium compound, only indium sulfate is shown in the figure.

In the present Embodiment, the result of the case using the one having a plating film thickness of 0.3μ m has been shown, however, in practice, also when the one having a film thickness of $0.1-1 \mu$ m was used, there was little difference in the result.

(Comparative example 3)

For comparison, an experiment was performed using non-plated copper plates in the same procedure as Embodiment 25. Experiment results are shown in Table 19 and Fig. 16.

Table 19

Hydrogen evolution rate with not plated copper plate			
Example No.	Electrolyte	Inhibitor (concentration: 100ppm)	Hydrogen evolution rate μ L/g/day
1	KOH	-	516.00
2	KOH	Indium sulfate	30.56
3	KOH	Indium sulfamate	28.73
4	KOH	Indium hydroxide	29.07
5	KOH	Lead oxide	24.15
6	KOH	Barium hydroxide	480.32
7	KOH	Carbon fluoride-POE	432.19

It is understood that in the case of no addition of the inhibitor and addition of barium hydroxide or the surfactant of the fluorocarbon-polyoxyethylene series, the hydrogen gas generation amount increases with respect to the number of test days exponentially. On the other hand, it is understood that in the case of addition of indium sulfate and lead oxide, the generation of hydrogen gas is suppressed. This is due to the fact that the compound of indium or lead has the function to coat the copper plate, but barium hydroxide and the surfactant have such a function a little.

(Embodiment 26)

Using a copper plate plated in the same manner as Example 25, the hydrogen gas generation in the case of the use of combination of inhibitors was measured using an electrolyte solution of the KOH type. Results are shown in Table 20.

Table 20

Hydrogen evolution rate with indium plated copper plate and inhibitor					
Experimental No.	Concentration of inhibitor ppm				Hydrogen evolution rate μ L/g/day
	Indium sulfate	Lead oxide	Indium hydroxide	POERA	
43	100	100	0	0	15.82
44	100	0	100	0	10.10
45	100	0	0	100	12.28
46	0	100	100	0	14.64
47	0	100	0	100	15.40
48	0	0	100	100	9.28
49	100	100	100	0	14.87
50	100	100	0	100	12.12
51	100	0	100	100	8.88
52	100	100	100	100	8.54

In the column for the inhibitor in the table, the concentration of each inhibitor added to the electrolyte solution with respect to the electrolyte solution is shown.

It is understood that the hydrogen gas generation is at a low level in the same manner as Example, and there is not too much bad effect even when a plurality of inhibitors are used. Especially, the hydrogen gas generation is less in the one in which barium hydroxide and polyoxyethylene alkylamide are simultaneously added.

5 Next, a battery was actually manufactured, and the effects of the plating on the collector or the inhibitor were investigated.

(Embodiment 27)

10 Into a positive electrode can were added a part of an electrolyte solution and 116 mg of a pellet (silver oxide content of 98 %) in which silver oxide was added and molded with a combined agent, and a separator of polyethylene and a separator of cellophane were placed. Next, a gasket of nylon was pushed and fitted into the positive electrode can, and filling with an impregnating agent, a gelling agent, and 30 mg of zinc powder, the remainder of the electrolyte solution optionally added with an inhibitor was added dropwise,
15 and then a negative electrode can was placed to perform sealing to manufacture 100 pieces for each kind of button type silver oxide batteries.

As the zinc powder was used one containing 130 ppm of bismuth, 500 ppm of indium and 30 ppm of aluminum manufactured by atomization.

20 The type and the film thickness of plating onto the negative electrode can, the type and the concentration of the added inhibitor, the closed circuit voltage and the self-discharge rate are shown in Tables 21 and 22.

Table 21. Characteristics of cell with KOH electrolyte

Experimental No.	Kind of plating	Concentration of inhibitor ppm				Self-discharge rate %		Closed circuit voltage V	
		Indium sulfate	Lead oxide	barium hydroxide	POERA	Fresh cell	A part- discharged cell	Fresh cell	A part- discharged cell
53	—	0	0	0	0	5.8	9.9	1.158	1.179
54	In	0	0	0	0	4.3	5.9	1.165	1.192
55	In	1000	0	0	0	2.7	2.7	1.169	1.198
56	In	0	1000	0	0	2.7	2.8	1.168	1.197
57	In	0	0	1000	0	2.5	4.9	1.185	1.198
58	In	0	0	0	1000	2.6	5.0	1.165	1.185
59	In	1000	1000	0	0	2.5	2.6	1.168	1.197
60	In	1000	0	1000	0	2.4	2.6	1.185	1.239
61	In	1000	0	0	1000	2.4	2.5	1.168	1.199
62	In	0	1000	0	1000	2.4	2.6	1.168	1.197
63	In	0	1000	1000	0	2.6	2.7	1.167	1.238
64	In	0	1000	1000	1000	2.4	2.4	1.188	1.239
65	In	1000	1000	1000	0	2.6	2.6	1.186	1.240
66	In	1000	1000	0	1000	2.5	2.6	1.170	1.200
67	In	1000	0	1000	1000	2.3	2.5	1.185	1.240
68	In	1000	1000	1000	1000	2.4	2.4	1.186	1.238
69	Sn	0	0	0	0	4.7	6.8	1.167	1.193
70	Sn	1000	0	0	0	2.8	2.7	1.168	1.198
71	Sn	0	1000	0	0	2.7	2.8	1.170	1.199
72	Sn	0	0	1000	0	2.6	5.0	1.187	1.199
73	Sn	0	0	0	1000	2.6	5.1	1.165	1.185
74	Sn	1000	1000	0	0	2.6	2.7	1.168	1.197
75	Sn	1000	0	1000	0	2.4	2.5	1.187	1.240
76	Sn	1000	0	0	1000	2.4	2.4	1.168	1.201
77	Sn	0	1000	0	1000	2.4	2.6	1.169	1.196
78	Sn	0	1000	1000	0	2.7	2.8	1.167	1.240
79	Sn	0	1000	1000	1000	2.6	2.5	1.189	1.240
80	Sn	1000	1000	1000	0	2.7	2.6	1.185	1.240
81	Sn	1000	1000	0	1000	2.5	2.6	1.172	1.200
82	Sn	1000	0	1000	1000	2.3	2.6	1.185	1.241
83	Sn	1000	1000	1000	1000	2.4	2.5	1.188	1.241
84	Zn·In	0	0	0	0	4.9	7.1	1.171	1.199
85	Zn·In	1000	0	0	0	2.6	2.7	1.167	1.197
86	Zn·In	0	1000	0	0	2.9	3.0	1.188	1.200
87	Zn·In	0	0	1000	0	2.6	5.0	1.165	1.186
88	Zn·In	0	0	0	1000	2.6	5.0	1.168	1.197
89	Zn·In	1000	1000	0	0	2.5	2.6	1.186	1.239
90	Zn·In	1000	0	1000	0	2.5	2.7	1.169	1.200
91	Zn·In	1000	0	0	1000	2.4	2.7	1.168	1.199
92	Zn·In	0	1000	0	1000	2.5	2.5	1.168	1.237
93	Zn·In	0	1000	1000	0	2.6	2.9	1.188	1.241
94	Zn·In	0	1000	1000	1000	2.5	2.5	1.187	1.241
95	Zn·In	1000	1000	1000	0	2.5	2.6	1.169	1.200
96	Zn·In	1000	1000	0	1000	2.7	2.6	1.187	1.240
97	Zn·In	1000	0	1000	1000	2.4	2.6	1.187	1.239
98	Zn·In	1000	1000	1000	1000	2.5	2.6	1.188	1.240

Table 22. Characteristics of cell with NaOH electrolyte

Experimental No.	Kind of plating	Concentration of inhibitor ppm				Self-discharge rate %		Closed circuit voltage V	
		Indium sulfate	Lead oxide	barium hydroxide	POERA	Fresh cell	A part-discharged cell	Fresh cell	A part-discharged cell
99	—	0	0	0	0	5.5	9.6	1.157	1.181
100	In	0	0	0	0	4.0	5.5	1.167	1.194
101	In	1000	0	0	0	2.3	2.4	1.169	1.199
102	In	0	1000	0	0	2.2	2.5	1.169	1.198
103	In	0	0	1000	0	2.2	4.7	1.185	1.199
104	In	0	0	0	1000	2.2	4.8	1.166	1.184
105	In	1000	1000	0	0	2.1	2.3	1.169	1.199
106	In	1000	0	1000	0	1.9	2.2	1.188	1.240
107	In	1000	0	0	1000	1.9	2.2	1.168	1.200
108	In	0	1000	0	1000	2.1	2.4	1.167	1.199
109	In	0	1000	1000	0	2.2	2.4	1.169	1.238
110	In	0	1000	1000	1000	1.8	1.9	1.189	1.240
111	In	1000	1000	1000	0	2.4	2.2	1.187	1.241
112	In	1000	1000	0	1000	2.2	2.1	1.171	1.202
113	In	1000	0	1000	1000	2.0	2.2	1.186	1.240
114	In	1000	1000	1000	1000	2.0	2.0	1.188	1.240
115	Sn	0	0	0	0	4.4	5.4	1.169	1.195
116	Sn	1000	0	0	0	2.5	2.2	1.169	1.200
117	Sn	0	1000	0	0	2.5	2.3	1.170	1.200
118	Sn	0	0	1000	0	2.4	4.7	1.189	1.200
119	Sn	0	0	0	1000	2.3	4.7	1.167	1.184
120	Sn	1000	1000	0	0	2.2	2.1	1.169	1.196
121	Sn	1000	0	1000	0	2.1	2.3	1.188	1.240
122	Sn	1000	0	0	1000	1.9	2.1	1.169	1.200
123	Sn	0	1000	0	1000	2.2	2.3	1.168	1.198
124	Sn	0	1000	1000	0	2.5	2.4	1.169	1.240
125	Sn	0	1000	1000	1000	2.4	2.2	1.190	1.240
126	Sn	1000	1000	1000	0	2.4	2.3	1.186	1.241
127	Sn	1000	1000	0	1000	2.3	2.4	1.174	1.199
128	Sn	1000	0	1000	1000	2.0	2.4	1.186	1.241
129	Sn	1000	1000	1000	1000	2.1	2.2	1.189	1.241
130	Zn·In	0	0	0	0	4.6	6.7	1.172	1.201
131	Zn·In	1000	0	0	0	2.3	2.4	1.169	1.198
132	Zn·In	0	1000	0	0	2.7	2.5	1.188	1.201
133	Zn·In	0	0	1000	0	2.4	4.8	1.168	1.187
134	Zn·In	0	0	0	1000	2.3	4.8	1.170	1.197
135	Zn·In	1000	1000	0	0	2.1	2.4	1.188	1.240
136	Zn·In	1000	0	1000	0	2.2	2.4	1.170	1.199
137	Zn·In	1000	0	0	1000	2.2	2.5	1.169	1.199
138	Zn·In	0	1000	0	1000	2.2	2.2	1.167	1.236
139	Zn·In	0	1000	1000	0	2.1	2.6	1.187	1.243
140	Zn·In	0	1000	1000	1000	2.3	2.0	1.187	1.242
141	Zn·In	1000	1000	1000	0	2.3	2.4	1.168	1.201
142	Zn·In	1000	1000	0	1000	2.2	2.4	1.189	1.241
143	Zn·In	1000	0	1000	1000	2.2	2.4	1.187	1.240
144	Zn·In	1000	1000	1000	1000	2.3	2.3	1.187	1.241

The inhibitor concentration in the table indicates a value with respect to the zinc weight. The measurement of the self-discharge rate was performed after maintaining at 60 °C for 20 days said to be equivalent to 1 year. The self-discharge rate after partial discharge was measured by being left at 60 °C for 20 days after 50 % depth discharge (partial discharge). The closed circuit voltage was measured at -10 °C before discharge and after partial discharge.

As compared with those of no addition (Example No. 54, 69 and 84 in Table 21 and Example No. 100, 115 and 130 in Table 22), those added with the inhibitor have small self-discharge rates. With respect to the

partial discharge, in the case of those added with one of the type for coating the collector and the zinc surface such as indium sulfate and lead monoxide, the self-discharge rate is improved. It is apparent that the closed circuit voltage is high in those added with barium hydroxide. It is understood from the table that when the inhibitor of the coating type and barium hydroxide are added, the self-discharge rate before discharge, the self-discharge rate after partial discharge, and the closed circuit voltage are improved. In the present experiment, a behavior that the closed circuit voltage after partial discharge becomes higher than that before discharge has been presented which is opposite to a case using mercurated zinc. The cause thereof is now under investigation.

The plating onto the negative electrode can was performed in the same manner as Example 25, and results for each film thickness of 0.3 μ m have been shown. There was little difference in the result even when those having a film thickness of 0.1-1 μ m were used.

(Embodiment 28)

Further, with a battery manufactured using an electrolyte of the KOH type and a negative electrode can applied with indium plating, the self-discharge rate corresponding to the one after 1 year and the self-discharge rate after partial discharge with respect to the inhibitor concentration were measured.

Results are shown in Figs. 17 and 18. Provided that the practical range is not more than about 3 % of the self-discharge rate, according to Fig. 17, it has been found to be effective for indium sulfate of 50-5000 ppm, for lead monoxide of 20-5000 ppm, for barium hydroxide of not less than 50 ppm, and for polyoxyethylene alkylamide of not less than 5 ppm as the concentration with respect to zinc.

With respect to the self-discharge rate after 50 % depth discharge, according to Fig. 18, it was effective for indium sulfate of 50-5000 ppm, and for lead monoxide of 20-5000 ppm. After partial discharge, the effect is not so much for barium hydroxide and polyoxyethylene alkylamide. In order to lower the self-discharge rate after partial discharge, it has been found to be effective to combine and use indium sulfate or lead monoxide and barium hydroxide-polyoxyethylene alkylamide. However, it is desirable that, in the combined use polyoxyethylene alkylamide is not more than 1000 ppm. The result has been shown in the case of indium sulfate for indium compounds, however, other compounds were also effective was in substantially the same concentration range. In addition, even when an electrolyte solution of the NaOH type or other negative electrode can plating were used, the concentration range of the inhibitor exhibiting the effect was substantially the same.

(Embodiment 29)

Trial production of batteries was performed in the same manner as Example 27 by changing the zinc composition. Using an electrolyte solution of the KOH type and plating of indium onto a negative electrode can, and as the inhibitor, each 1000 ppm of indium sulfate, lead monoxide, barium hydroxide and polyoxyethylene alkylamide with respect to zinc was added. Results are shown in Table 23.

Table 23. Characteristics of cell with KOH electrolyte

Experi- mental No.	Concentration of metal added to zinc ppm					Self-discharge rate %		Closed circuit voltage V	
	Pb	Bi	In	Al	Ca	Fresh cell	A part- discharged cell	Fresh cell	A part- discharged cell
145	500	-	-	-	-	2.9	2.8	1.186	1.200
146	500	50	-	-	-	2.8	2.8	1.187	1.235
147	500	-	-	250	-	2.8	2.7	1.186	1.235
148	500	500	500	-	-	2.3	2.5	1.186	1.234
149	10	100	515	30	-	2.7	2.7	1.188	1.222
150	23	240	-	-	-	2.9	2.9	1.189	1.200
151	28	240	1780	-	-	2.5	2.6	1.189	1.240
152	-	140	480	-	-	2.4	2.6	1.185	1.239
153	-	130	500	30	-	2.4	2.4	1.186	1.238
154	-	130	500	-	50	2.3	2.4	1.188	1.236

It is apparent that the self-discharge rate is included in a practical range of not more than 3 % in the compositions of additives defined in claims. In order to lower the self-discharge rate, it is desirable to form an alloy of several species.

In Examples 30-31, results of evaluation mainly for combinations with gelling agents are shown.

5
(Embodiment 30)

Before practically manufacturing batteries, the effect of the present invention was confirmed in a form of a hydrogen gas generation test. The test was performed using a combination of zinc powder, a gelling agent of the present invention, barium hydroxide and a copper plate of the same material as that of the collector. Into a specially prepared test tube having a volume of 25 ml graduated to know the gas generation amount was added 2 g of zinc powder containing 500 ppm of each of bismuth, indium and lead manufactured by the atomization method, and an electrolyte solution to be tested was added thereto and heated to 60 °C, and the volume of generated hydrogen gas was measured for 7 days. The gelling agent was added by 1.5 % with respect to zinc. Using 2 g of zinc powder and 5 copper pieces as the same material as the collector having a surface area of 0.6 cm² and a thickness of 0.1 mm together so as to provide the same ratio of the negative electrode zinc weight to the negative electrode collector surface area of the battery, the gas generation was measured in the same manner. The number of repetition in the test was 6, and an average value thereof was used as a result.

20 The electrolyte solution was prepared by providing a base solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide type, or a base solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide type, and barium hydroxide was added thereto. Barium hydroxide was made by Wako Pure Chemical Industries., Ltd., CMC was No. 1260 and No. 1380 made by Daicel Chemical Industries, Ltd., and PAS was a reagent made by Wako Pure Chemical Industries., Ltd. and Rheogic 250H made by Nihon Pure Chemicals Co., Ltd. The added amount of barium hydroxide was 0-50000 ppm with respect to the electrolyte solution. Results are shown in the hydrogen generation amount in Table 24.

Table 24. Hydrogen evolution rate

Experi- mental No.	Kind of electro- lyte	Gelled agent	Copper pieces	Amount of added barium hydroxide ppm	Hydrogen evolution rate $\mu\text{l/g/day}$
1	KOH	CMC 1260	none	0	35.71
2	KOH	CMC 1380	none	0	30.43
3	KOH	PAS WAKO	none	0	22.12
4	KOH	PAS 250H	none	0	13.58
5	KOH	CMC 1260	none	1000	24.73
6	KOH	CMC 1380	none	1000	19.39
7	KOH	PAS WAKO	none	1000	8.65
8	KOH	PAS 250H	none	1000	6.82
9	KOH	CMC 1260	include	1000	215.31
10	KOH	CMC 1380	include	1000	189.56
11	KOH	PAS WAKO	include	1000	99.49
12	KOH	PAS 250H	include	1000	81.72
13	KOH	none	none	50000	2.02
14	KOH	none	none	10000	1.99
15	KOH	none	none	5000	2.01
16	KOH	none	none	1000	2.10
17	KOH	none	none	500	2.24
18	KOH	none	none	100	2.55
19	KOH	none	none	0	3.19
20	KOH	none	include	50000	238.42
21	KOH	none	include	10000	31.54
22	KOH	none	include	1000	46.37
23	NaOH	none	include	100	84.25
24	NaOH	none	include	0	516.63
25	NaOH	none	none	50000	1.90
26	NaOH	none	none	10000	1.95
27	NaOH	none	none	5000	2.00
28	NaOH	none	none	1000	2.02
29	NaOH	none	none	500	2.12
30	NaOH	none	none	100	2.31
31	NaOH	none	none	0	3.46
32	NaOH	none	include	50000	25.13
33	NaOH	none	include	10000	27.58
34	NaOH	none	include	1000	31.49
35	NaOH	none	include	100	95.61
36	NaOH	none	include	0	252.89

The unit is $\mu\text{l/g/day}$. According to the result in Table 24, it is understood that one added with barium hydroxide has a small hydrogen generation amount irrespective to comparison in the gelling agent type, the zinc powder simple substance and the zinc powder + copper piece, and the electrolyte solution type, and the gas generation is small in PAS being a cross-linked type acrylic water-soluble resin with respect to the gelling agent type.

It is apparent that the combination of PAS as a cross-linked type acrylic water-soluble resin and barium hydroxide is especially good. When the adding amount of barium hydroxide was not less than 500 ppm, there was supersaturation, so that the electrolyte solution in a turbid state was used. In addition, barium hydroxide was added to the electrolyte solution in Example 30, however, the same effect was obtained even when the whole amount was added to powdery zinc.

(Embodiment 31)

A button type silver oxide battery of the SR621 size was prepared. As a negative electrode can the one in which a collector was applied with tin plating of $0.1 \mu\text{m}$ was used. The zinc was the one used in Example 30, Rheogic 250H as a cross-linked type acrylic water-soluble resin for a gelling agent, and a potassium hydroxide 30 % by weight solution or a sodium hydroxide 25 % by weight solution added with zinc oxide up to approximate saturation respectively for an electrolyte solution. The adding amount of the gelling agent was 1.5 % with respect to zinc, and the adding amount of barium hydroxide was 0-50000 ppm with respect to zinc. As Conventional example, a battery was manufactured in which CMC 1260# was used for the gelling agent with no barium hydroxide added. Evaluation results of electric characteristics are shown in the discharge index in Table 25.

Table 25

Chacteristics of cell				
Experimental No.	Kind of electrolyte	Amount of added barium hydroxide ppm	Comparison of discharge capacity	Hydrogen evolution rate $\mu\text{ l/g/day}$
1	KOH	50000	97	0.208
2	KOH	10000	100	0.205
3	KOH	1000	103	0.227
4	KOH	100	1023	0.708
5	KOH	10	102	1.693
6	KOH	0	101	1.812
Conventional	KOH	0	100	2.513
7	NaOH	50000	98	0.192
8	NaOH	10000	101	0.185
9	NaOH	1000	102	0.183
10	NaOH	100	102	0.228
11	NaOH	10	101	0.342
12	NaOH	0	101	0.549
Conventional	NaOH	0	100	0.712

The electric characteristics were measured by the direct current method with a load resistance of 200Ω when the electrolyte solution was the potassium hydroxide type or the pulse method with a load resistance of $2 \text{ k} \Omega$ when it was the sodium hydroxide type. In either case, Conventional example was supposed to have a discharge index of 100. As being apparent also from the result, one in which the gelling agent of the present invention and the cross-linked type acrylic water-soluble resin were used and barium hydroxide was added had a good discharge performance as compared with Conventional example.

Using a gasket of polypropylene specially prepared for allowing hydrogen to penetrate, a button type silver oxide battery was manufactured in the same manner. As a gelling agent, 1.5 % of Rheogic 250H being a cross-linked type acrylic water-soluble resin was used with respect to zinc. Barium hydroxide had been tested with a concentration of 10 ppm to 5 % with respect to a weight of zinc. Ten pieced of the manufactured silver oxide batteries were placed in a vessel made of glass filled with liquid paraffin in a high temperature tank, a collecting tube having graduations at an upper portion was attached, and the amount of generated hydrogen was measured. This state was maintained at 60°C for 20 days, and the hydrogen gas generation amount after 20 days was investigated. Evaluation results are shown in the hydrogen gas generation amount in Table 25, and a result of a case using a potassium hydroxide 30 % by weight solution as the electrolyte solution is shown in Fig. 19. According to the figure, it is apparent that barium hydroxide works effectively at 100 ppm to 1 %.

The gasket made of polypropylene was returned to one made of nylon, and a button type silver oxide battery was manufactured in a range of barium hydroxide concentration of 100 ppm to 1 % which had less gas generation. Ten pieces of the manufactured batteries were placed in a vessel made of glass filled with liquid paraffin in a high temperature tank in the same manner, and a collecting tube for generating hydrogen gas was attached to an upper portion. Hydrogen gas generation, expansion of the battery can and liquid leakage were not observed after 20 days at 60°C .

The hydrogen gas generation amount in the actual battery has a value smaller than that of the hydrogen gas generation amount in the experiment in Example 30. For this fact, it is considered that the copper piece used in Example 30 is different from a structure of an actual collector, a part of generated hydrogen gas has been consumed by the reduction of silver oxide and the like. The method in Example 30 is different in the amount order of hydrogen gas generation, but it is sufficient as a method for observing substituted characteristics for predicting hydrogen generation in an actual battery.

In accordance with Examples 32-35, the effect of a case in which the collector is provided with a zinc alloy layer and evaluation results of a case in which the alkaline battery of the present invention is used for a clock or watch are shown.

Before actually investigating characteristics using batteries, a specially prepared test tube was used to investigate what amount of hydrogen gas was generated from a plated copper plate (the same material as a collector). The method therefor will be described hereinafter.

Into a specially prepared test tube having a volume of 25 ml graduated to know the gas generation amount were placed beforehand 2 g of zinc powder containing 500 ppm of each of bismuth, indium and lead manufactured by the atomization method, and a copper piece being the same material as the collector and having an area of 0.6 cm^2 and a thickness of 0.1 mm being applied with plating of the present invention, an electrolyte solution was added thereto and heated to 60°C , and the volume of generated hydrogen gas was measured for 7 days. The number of repetition of the test was 10, and an average value thereof was used as a result. As the electrolyte solution was used a solution in which potassium hydroxide was 30 % by weight and zinc oxide was added up to approximate saturation in the case of the potassium hydroxide type, and a solution in which sodium hydroxide was 25 % by weight and zinc oxide was added up to approximate saturation in the case of the sodium hydroxide type.

(Embodiment 32)

The hydrogen generation from zinc and a copper piece coated with the electric plating and the substitution plating was measured. Four kinds of plating treatments onto the copper piece were performed. Conditions are shown hereinafter.

= Contact plating (zinc-indium)

method: contact plating: A copper piece and zinc were placed into the following plating bath heated to 60°C and left for 1 hour.

plating bath: KOH 30 %, ZnO saturation, indium sulfate 0-3000 ppm

temperature: 60°C

plating film thickness: Adjustment was made depending on time. The thickness was determined from weights before and after the plating.

= Contact plating (zinc-lead)

method: contact plating: A copper piece and zinc were placed into the following plating bath heated to 60°C and left for 1 hour.

plating bath: KOH 30 %, ZnO saturation, lead oxide 0-3000 ppm

temperature: 60°C

plating film thickness: Adjustment was made depending on time. The thickness was determined from weights before and after the plating.

= Electric plating (cyanide type)

method: A copper piece was clamped in a net made of copper to allow a current to flow, and plating was performed. The copper piece was suitably moved so as not to make a portion not subjected to plating.

plating bath: zinc cyanide 70 g/l, sodium cyanide 50 g/l, sodium hydroxide 100 g/l, indium cyanide 0-3000 ppm

temperature: 25°C

current density: 5 A/dm^2

plating film thickness: Adjustment was made depending on time. The thickness was determined from weights before and after the plating.

= Electric plating (acidic)

method: A copper piece was clamped in a net made of copper to allow a current to flow, and plating was performed. The copper piece was suitably moved so as not to make a portion not subjected to plating.

5 plating bath: zinc sulfate 240 g/l, sodium sulfate 30 g/l, sodium acetate 15 g/l, indium sulfate 0-3000 ppm

temperature: 25 °C

current density: 2 A/dm²

10 plating film thickness: Adjustment was made depending on time. The thickness was determined from weights before and after the plating.

A copper piece subjected to each treatment, zinc powder and an electrolyte solution were added in a specially prepared test tube, and the amount of hydrogen gas generated therefrom was investigated. It is supposed that the larger the hydrogen gas generation, the larger is the self-discharge in an actual battery.

Table 26

Hydrogen evolution rate in KOH electrolyte							
Experimental No.	Condition of plating		Hydrogen evolution rate (μ l/g/day) vs concentration of additive in plating solution (ppm)				
	Kind of plating	Additive	0	10	100	1000	3000
1	Contact plating	Indium sulfate	516.92	14.29	10.71	12.14	8.57
2	Contact plating	PbO	↑	13.28	12.96	96.34	120.36
3	Electroplating	Indium cyanide	↑	15.47	13.22	10.25	10.81
4	Electroplating	Indium sulfate	↑	18.24	15.26	15.13	7.41

Table 27

Hydrogen evolution rate in NaOH electrolyte							
Experimental No.	Condition of plating		Hydrogen evolution rate (μ l/g/day) vs concentration of additive in plating solution (ppm)				
	Kind of plating	Additive	0	10	100	1000	3000
5	Contact plating	Indium sulfate	480.23	13.00	10.17	11.53	7.11
6	Contact plating	PbO	↑	12.75	11.28	90.56	102.31
7	Electroplating	Indium cyanide	↑	14.54	12.82	9.43	9.84
8	Electroplating	Indium sulfate	↑	18.06	14.19	14.98	6.45

In Table 26, the hydrogen generation amount in the KOH type electrolyte solution with respect to the copper piece plated with each concentration of lead oxide is shown. In Table 27, the result in the NaOH type electrolyte solution is shown.

According to Tables 26 and 27, it is understood that the hydrogen gas generation can be greatly suppressed as by adding a little amount of indium compound and lead oxide of about 10 ppm into the plating solution, and forming an alloy of the plating film as compared with no addition. However, in the case of addition of lead oxide in a high concentration, needle-like lead becomes deposited on the plating film, and the hydrogen gas generation is deteriorated.

When the copper piece subjected to the substitution plating was quantitatively measured by the Auger spectroscopic analysis, it was found that about 0.3 % of indium existed in the plating film in the case of adding indium sulfate in an amount of 1000 ppm, and about 0.1 % of indium existed in the case of adding an amount of 100 ppm. In the case of the plating film manufactured by adding indium sulfate to the plating solution by not more than 100 ppm, it was impossible to measure indium as being not more than the

detection limit. However, it is considered that the hydrogen gas generation suppressing effect is obtained by the inclusion of an extremely minute amount of indium into the plating film. For other kinds of plating, it is considered that, in the same manner the hydrogen gas generation suppressing effect is obtained because indium and lead are deposited in zinc, to lower the hydrogen overvoltage of zinc.

Table 28. Hydrogen evolution rate in KOH electrolyte

Experi- mental No.	Condition of plating		Hydrogen evolution rate ($\mu\text{l/g/day}$) vs plating thickness (μm)					
	Kind of plating	Additive	0	0.015	0.074	0.082	0.152	0.222
9	Contact plating	Indium sulfate	516.9	11.79	7.50	13.21	8.93	8.75

Table 29. Hydrogen evolution rate in NaOH electrolyte

Experi- mental No.	Condition of plating		Hydrogen evolution rate ($\mu\text{l/g/day}$) vs plating thickness (μm)					
	Kind of plating	Additive	0	0.015	0.074	0.082	0.152	0.222
10	Contact plating	Indium sulfate	480.2	12.36	6.23	6.02	7.71	7.11

In Tables 28 and 29, the hydrogen gas generation amount with respect to the film thickness is shown in the substitution plating added with 1000 ppm of indium sulfate. According to Tables 28 and 29, it is apparent that the effect is provided from 0.015 μm as being fairly thin.

(Embodiment 33)

Next, trial production of a battery was actually performed to confirm the effect on the negative electrode can.

Into a positive electrode can were added a part of an electrolyte solution and placing therein 116 mg of a pellet (silver oxide content of 98 %) made by adding silver oxide being and molded with a combined agent, a separator of polyethylene and a separator of cellophane. Next, a gasket of nylon was pushed and fitted into the positive electrode can, an impregnating agent, a gelling agent, and 30 mg of zinc powder were added, the remainder of the electrolyte solution optionally added with an inhibitor was added dropwise, then a negative electrode can was placed and sealed to manufacture 100 pieces for each kind of button type silver oxide batteries.

As the zinc powder was used one containing 500 ppm of bismuth, 500 ppm of indium and 500 ppm of lead manufactured by atomization. The plating onto the negative electrode can was performed in the same manner as Example 32.

The self-discharge rate with respect to the type of the plating onto the negative electrode can is shown in Tables 30 and 31. The measurement of the self-discharge rate was performed after maintaining 60 °C for 20 days as said to be equivalent to 1 year.

Table 30

Self-discharge rate of cell with KOH electrolyte			
Experimental No.	Condition of plating		Self-discharge rate after 1 year storage %
	Kind of plating	Additive	
11	none	-	5.8
12	Contact plating	Indium sulfate	2.5
13	Contact plating	PbO	2.8
14	Electroplating	Indium cyanide	2.4
15	Electroplating	Indium sulfate	2.4

Table 31

Self-discharge rate of cell with NaOH electrolyte			
Experimental No.	Condition of plating		Self-discharge rate after 1 year storage %
	Kind of plating	Additive	
16	none	-	5.5
17	Contact plating	Indium sulfate	2.2
18	Contact plating	PbO	2.5
19	Electroplating	Indium cyanide	2.1
20	Electroplating	Indium sulfate	2.2

According to Tables 30 and 31, it is understood that the self-discharge can be suppressed by using the plated negative electrode can of the present invention as compared with the negative electrode can not subjected to plating.

In this case, results are shown in which the adding amount of the indium Compound into the plating solution was 1000 ppm, the adding amount of lead oxide was 100 ppm, and each film thickness was 0.1 μ m. There was little difference in the result even when those having a film thickness of 0.01-3 μ m were used. In addition, equivalent effects were obtained in ranges in which the amount of the indium compound added into the plating solution was 10-3000 ppm, and the amount of lead oxide added was 10-100 ppm. When the same test was performed for potassium stannate and sodium stannate, equivalent effects were obtained in a range of the adding amount of 10-3000 ppm. In addition, equivalent effects were obtained even when plating solutions mixed and added with indium compounds, lead oxide, potassium stannate and sodium stannate in these concentration ranges have been used.

(Embodiment 34)

A test was performed for the impurity shielding effect of the plating film.

Plating was performed for a negative electrode can having the SR521 size in which iron type foreign matters had adhered to the collector surface during press processing causing deficiency in gas generation. The deficiency in gas generation may become a cause of expansion and explosion when a battery is made, which can be confirmed by instant gas generation after adding an electrolyte solution and several zinc powder particles of about 100 μ m to a concave surface at the copper side being the collector of the negative electrode can. This is due to the fact that zinc serves as an anode and iron having a lower hydrogen overvoltage serves as a cathode to generate hydrogen. The plating was performed by the substitution plating using a barrel. Conditions are described hereinafter.

= Contact plating (zinc-indium)

method: contact plating: Using a barrel, a negative electrode can and a zinc piece were added into the following plating solution heated to 60 °C to perform plating.

5 negative electrode can; SR521 size, about 1000 pieces

zinc piece: 3 x 3 x 0.1, about 1500 sheets

plating solution: KOH 30 %, ZnO saturation, indium sulfate 0-300 ppm

temperature: 60 °C

time: 5, 15 and 30 minutes

10 plating film thickness: The thickness was measured from weights of a copper plated added as a dummy before and after the plating.

The film thickness against time was 0.041 μ m for 5 minutes, 0.090 μ m for 15 minutes, and 0.148 μ m for 30 minutes. 100 pieces were picked out respectively, and the confirmation of gas generation was performed. As a result, the gas generation was found in 78 pieces out of 100 pieces in the case of those subjected to plating for 5 minutes. No gas generation was found in those subjected to plating for 15 and 30 minutes. According to this fact, it has been found that a plating thickness of about 0.1 μ m has shielding effect for impurities.

(Embodiment 35)

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Next, results are shown for a case in which the shaping into a negative electrode can was performed after plating onto a hoop material instead of the direct negative electrode can plating as done in Example 33. In this case, although there is a merit that the plating can be made thick by the electric plating, however, there is a problem that deficiency is apt to be caused in a soft plating film by mechanical processing. Plating conditions are shown hereinafter.

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* Electric plating (cyanide type)

method: The nickel side of a hoop material clad of three-layers of nickel, stainless and copper was subjected to masking, and plating was performed onto the copper side.

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plating bath: zinc cyanide 70 g/l, sodium cyanide 50 g/l, sodium hydroxide 100 g/l, indium cyanide 1000 ppm

bath temperature: 25 °C

Current density: 5 A/dm²

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plating film thickness: Adjustment was made depending on time. The thickness (adjustment of the feed speed of the hoop material) was determined by embedding in a resin and taking a picture using an optical microscope.

Those having a film thickness of 1, 2, 3, 4, 7 or 10 μ m were manufactured. Negative electrode cans of 100 pieces were manufactured using each hoop material plated with a zinc alloy of each film thickness. The number of negative electrode cans in which the copper surface became exposed by processing was investigated. Results are shown in Table 7.

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According to the result, it is apparent that the copper surface ceases to be exposed in the case of not less than 3 μ m.

The negative electrode cans of each film thickness were used to perform trial production of 100 pieces of batteries respectively with an NaOH type electrolyte solution in the same manner as Example 33.

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Table 32

Number of anode cap with generated pinhole and self-discharge rate							
Plating thickness (μ m)	1	2	3	4	7	10	Cell using amalgamated zinc
Number of anode cap with generated pinhole	8	3	0	0	0	0	
Initial capacity (mAh)	23.2	23.3	23.4	23.6	24.1	24.8	23.1
Capacity after 2 years storage (mAh)	21.2	21.7	22.0	22.7	23.0	23.8	22.6
Self-discharge rate (%/year)	4.4	3.5	2.9	1.9	2.1	2.0	1.1

The initial capacity and the capacity corresponding to the one after 2 years (after storage at 60 °C for 40 days) are shown in Table 32. A result of a case where mercurated zinc was used, is also described at the right end in the table for the purpose of comparison. According to Table 32, the plating of not more than 2 μ m has a capacity smaller than that of the case in which direct plating was performed to the negative electrode can in Example 33, due to deficiency in the plating film by processing. When the film thickness is 4 μ m, the capacity after 2 years becomes approximately the same as the one in which mercurated zinc is used. For example, when the battery is incorporated into a clock or watch, the battery service life is about 2 years. In order to expect a capacity equivalent to or not less than that of a battery using mercurated zinc after 2 years, it is necessary to plate the zinc alloy by not less than 4 μ m according to Table 32. In the design of a battery, it is available that the zinc alloy plating thickness may be not less than 3 μ m so as to have a thickness with which an aimed capacity is obtained in 2 years.

(Embodiment 36)

A conventional battery in which zinc having a mercuration ratio of 10 % was used and a battery in which a 4 μ m zinc-indium alloy was applied to a negative electrode can of the present invention by the same method as Example 35 using mercuryless zinc were actually installed in wristwatches, and a carrying test was performed to measure the number of days until the watch stopped. The test was done for each 10 pieces.

Table 33

Cell life by watch		
	Cell using amalgamated zinc	Cell using zinc without mercury and plated anode cap (plating thickness: 4 μ m)
Cell life(days)	751.1	758.3
σ	14.8	23.6

Results are shown in Table 33. The number of days until the watch stopped was substantially the same, and coming off with the capacity not being lowered even in the case of no mercury.

As shown in Example 35, it is apparent that when using as a standard a film thickness, in which the capacity after 2 years in the acceleration test becomes the same as that of the battery using mercurated zinc, it is possible to obtain a capacity equivalent to the one in which mercurated zinc is used also when used in an actual clock or watch.

Namely, in the design of a battery, the zinc alloy plating thickness may be made to be not less than 3 μ m so as to provide a thickness with which an aimed capacity is obtained in 2 years. For example, provided that the weight of zinc in the battery using mercurated zinc of an aimed capacity (excluding an amount of mercury) is W0, the self-discharge rate / year is SD0, the weight of zinc in the battery using mercuryless zinc to be designed is W0 (the same as mercurated zinc), the self-discharge rate is SD1, the area of the collector is S, and the specific gravity of zinc is 7.13, the film thickness D1 of the zinc alloy may be as follows:

$$D1 = (W0 \times (1 - 2 \times SD0) / (1 - 2 \times SD1) - W0) / 7.13 / S$$

In the present Examples 35 and 36, it has been described that the clad material is plated with the zinc-indium alloy of not less than 3 μ m, however, as shown in Examples 32 and 33, approximately the same effect is obtained provided that the zinc alloy is an alloy containing zinc as an essential element and containing one or more species selected from indium, lead and tin as a selective element. In addition, it is needless to say that the same effect is obtained even when the formation method of the zinc alloy layer is another method. For example, methods cladding a zinc alloy with one more layer, or using dry plating or flame coating and the like are applicable.

In addition, the effect of the invention is not deteriorated even when elements which raise the hydrogen overvoltage of the zinc alloy such as bismuth, gallium, aluminum, calcium and the like exist in the zinc alloy layer containing indium or lead. Depending on the way of addition, further self-discharge suppressing effects can be expected.

As also described in Examples hereinbefore, according to the present invention, it is possible to manufacture the alkaline battery in which the generation of hydrogen is suppressed without deteriorating battery characteristics. This is especially effective for coin type or button type alkaline batteries which require high performance but are sensitive to gas generation.

In addition, with respect to the zinc alloy plating onto the collector of the present invention, when it is used for minute discharge for clocks or watches or the like, it is possible to achieve a capacity which is equivalent to or not less than those of batteries in which mercurated zinc is used. Further, when clocks or watches carrying the alkaline batteries of the present invention are discarded, the environment is never polluted because mercury as an environmental pollution substance is not contained. When a applied product carrying the alkaline batteries are discarded, the environment is never polluted because mercury as an environmental pollution substance is not contained.

Claims

1. An alkaline battery comprising:
mercuryless zinc powder as a negative electrode active material; and
an electrolyte solution, the electrolyte Solution being added with any one species selected from indium sulfate, indium sulfamate and indium chloride.
2. An alkaline battery comprising:
mercuryless zinc powder as a negative electrode active material, the negative electrode active material being added with any one species selected from indium sulfate, indium sulfamate and indium chloride.
3. An alkaline battery comprising:
mercuryless zinc powder as a negative electrode active material; and
an electrolyte solution, the electrolyte solution being added with an indium compound and a complexing agent.
4. An alkaline battery according to claim 3 wherein said complexing agent is any one species selected from tartarate, EDTA and cyanate.
5. A method for producing alkaline batteries according to claim 3 comprising:
a step for dissolving a complexing agent in the electrolyte solution prior to the indium compound.
6. An alkaline battery comprising:
mercuryless zinc powder as a negative electrode active material, the negative electrode active material being added with an indium compound and a complexing agent.
7. An alkaline battery according to claim 6 wherein said complexing agent is any one species selected from tartarate, EDTA and cyanate.
8. An alkaline battery comprising:
mercuryless zinc powder as a negative electrode active material; and
an electrolyte solution, the electrolyte solution being added with two or more speices of compounds

selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

9. An alkaline battery according to claim 8 wherein said indium compound is indium sulfate, indium sulfamate, indium chloride and indium hydroxide.

10. An alkaline battery according to claim 8 wherein said tetravalent tin compound is a stannate such as potassium stannate, sodium stannate or the like and stannic oxide.

11. An alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material, the negative electrode active material being added with two or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

12. An alkaline battery according to claim 11 wherein said indium compound is indium sulfate, indium sulfamate, indium chloride and indium hydroxide.

13. An alkaline battery according to claim 11 wherein said tetravalent tin compound is a stannate such as potassium stannate, sodium stannate or the like and stannic oxide.

14. An alkaline battery comprising:

mercuryless zinc powder containing indium as a negative electrode active material; and an electrolyte solution, the electrolyte solution being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

15. An alkaline battery comprising:

mercuryless zinc powder containing indium as a negative electrode active material, the negative electrode active material being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

16. An alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material, a concentration of iron included in the mercuryless zinc powder being not more than 4 ppm with respect to zinc; and an electrolyte solution, the electrolyte solution being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

17. An alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material, a concentration of iron included in the mercuryless zinc powder being not more than 4 ppm with respect to zinc, the negative electrode active material being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

18. A silver oxide battery comprising:

mercuryless zinc as a negative electrode active material, the mercuryless zinc containing at least one species of metal selected from the group consisting of 0.001-0.5 % by weight of gallium, 0.005-0.5 % by weight of indium, 0.0001-0.5 % by weight of lead, 0.005-0.5 % by weight of bismuth, 0.001-0.5 % by weight of aluminum, and 0.001-0.5 % by weight of calcium, and the balance comprising zinc and inevitable impurities;

an amount of water in the battery being 0.31-0.57 mg per a weight of 1 mg of the mercuryless zinc; and

an electrolyte being potassium hydroxide.

19. A silver oxide battery comprising:

mercuryless zinc as a negative electrode active material, the mercuryless zinc containing at least one species of metal selected from the group consisting of 0.001-0.5 % by weight of gallium, 0.005-0.5 % by weight of indium, 0.0001-0.5 % by weight of lead, 0.005-0.5 % by weight of bismuth, 0.001-0.5 % by weight of aluminum, and 0.001-0.5 % by weight of calcium, and the balance comprising zinc and inevitable impurities;

an amount of water in the battery being 0.32-0.59 mg per a weight of 1 mg of the mercuryless zinc;

and
an electrolyte being sodium hydroxide.

20. An alkaline battery comprising:
 - mercuryless zinc powder containing a gelling agent as a negative electrode active material;
 - an outermost layer of a collector comprising a layer of zinc or metal having a hydrogen overvoltage higher than that of zinc; and
 - an electrolyte solution, the electrolyte solution being added with one or more species of materials selected from an indium compound, lead oxide, hydroxide of alkaline earth metal, a surfactant having polyethylene oxide.
21. The alkaline battery according to claim 20 wherein said zinc gelling agent is a cross-linked type polyacrylic water-absorbing polymer alone or a combination thereof with other gelling agents.
22. The alkaline battery according to claim 20 wherein said mercuryless zinc powder is mercuryless zinc powder containing at least one species of the group of 0.001-0.5 % by weight of gallium, 0.005-0.5 % by weight of indium, 0.0001-0.5 % by weight of lead, 0.005-0.5 % by weight of bismuth, 0.001-0.5 % by weight of aluminum, and 0.001-0.5 % by weight of calcium.
23. The alkaline battery according to claim 20 wherein said layer of zinc or metal having a hydrogen overvoltage higher than that of zinc is one or more species of metal or alloy selected from zinc, indium, tin and lead.
24. The alkaline battery according to claim 20 wherein said indium compound is indium sulfate, indium sulfamate, indium chloride and indium hydroxide.
25. The alkaline battery according to claim 20 wherein said hydroxide of alkaline earth metal is barium hydroxide.
26. The alkaline battery according to claim 20 wherein said surfactant having polyethylene oxide is added with one or more species of materials selected from a surfactant having fluorocarbon at the hydrophobic portion and polyethylene oxide at the hydrophilic portion, or polyoxyethylene alkylamide.
27. An alkaline battery comprising:
 - mercuryless zinc powder containing a gelling agent as a negative electrode active material, the negative electrode active material being added with one or more species of materials selected from an indium compound, lead oxide, hydroxide of alkaline earth metal, a surfactant having polyethylene oxide; and
 - an outermost layer of a collector comprising a layer of zinc or metal having a hydrogen overvoltage higher than that of zinc.
28. The alkaline battery according to claim 27 wherein said zinc gelling agent is a cross-linked type polyacrylic water-absorbing polymer alone or a combination thereof with other gelling agents.
29. The alkaline battery according to claim 27 wherein said mercuryless zinc powder is mercuryless zinc powder containing at least one species of the group of 0.001-0.5 % by weight of gallium, 0.005-0.5 % by weight of indium, 0.0001-0.5 % by weight of lead, 0.005-0.5 % by weight of bismuth, 0.001-0.5 % by weight of aluminum, and 0.001-0.5 % by weight of calcium.
30. The alkaline battery according to claim 27 wherein said layer of zinc or metal having a hydrogen overvoltage higher than that of zinc is one or more species of metal or alloy selected from zinc, indium, tin and lead.
31. The alkaline battery according to claim 27 wherein said indium compound is indium sulfate, indium sulfamate, indium chloride or indium hydroxide.
32. The alkaline battery according to claim 27 wherein said hydroxide of alkaline earth metal is barium hydroxide.

33. The alkaline battery according to claim 28 characterized in that said surfactant having polyethylene oxide is added with one or more species of materials selected from a surfactant having fluorocarbon at the hydrophobic portion and polyethylene oxide at the hydrophilic portion, or polyoxyethylene alkylamide.

34. An alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material; and
an outermost layer of a collector comprising an alloy layer containing zinc as an essential element, and containing one or more species selected from indium, lead and tin as a selective element.

35. The alkaline battery according to claim 34 wherein said alloy layer is formed by plating said outermost layer of said collector.

36. An alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material; and
an outermost layer of a collector comprising an alloy layer containing zinc as an essential element, and containing one or more species selected from indium, lead and tin as a selective element, the alloy layer being not less than 3 μ m.

37. An applied product using an alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material; and
an outermost layer of a collector comprising an alloy layer containing zinc as an essential element, and containing one or more species selected from indium and lead as a selective element, the alloy layer being not less than 3 μ m.

38. An applied product using an alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material; and
an electrolyte solution, the electrolyte solution being added with any one species selected from indium sulfate, indium sulfamate and indium chloride.

39. An applied product using an alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material, the negative electrode active material being added with any one species selected from indium sulfate, indium sulfamate and indium chloride.

40. An applied product using an alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material; and
an electrolyte solution, the electrolyte solution being added with an indium compound and a complexing agent.

41. An applied product using an alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material, the negative electrode active material being added with an indium compound and a complexing agent.

42. An applied product using an alkaline battery comprising:

mercuryless zinc powder as a negative electrode active material; and
an electrolyte solution, the electrolyte solution being added with two or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

43. An applied product using an alkaline battery comprising:

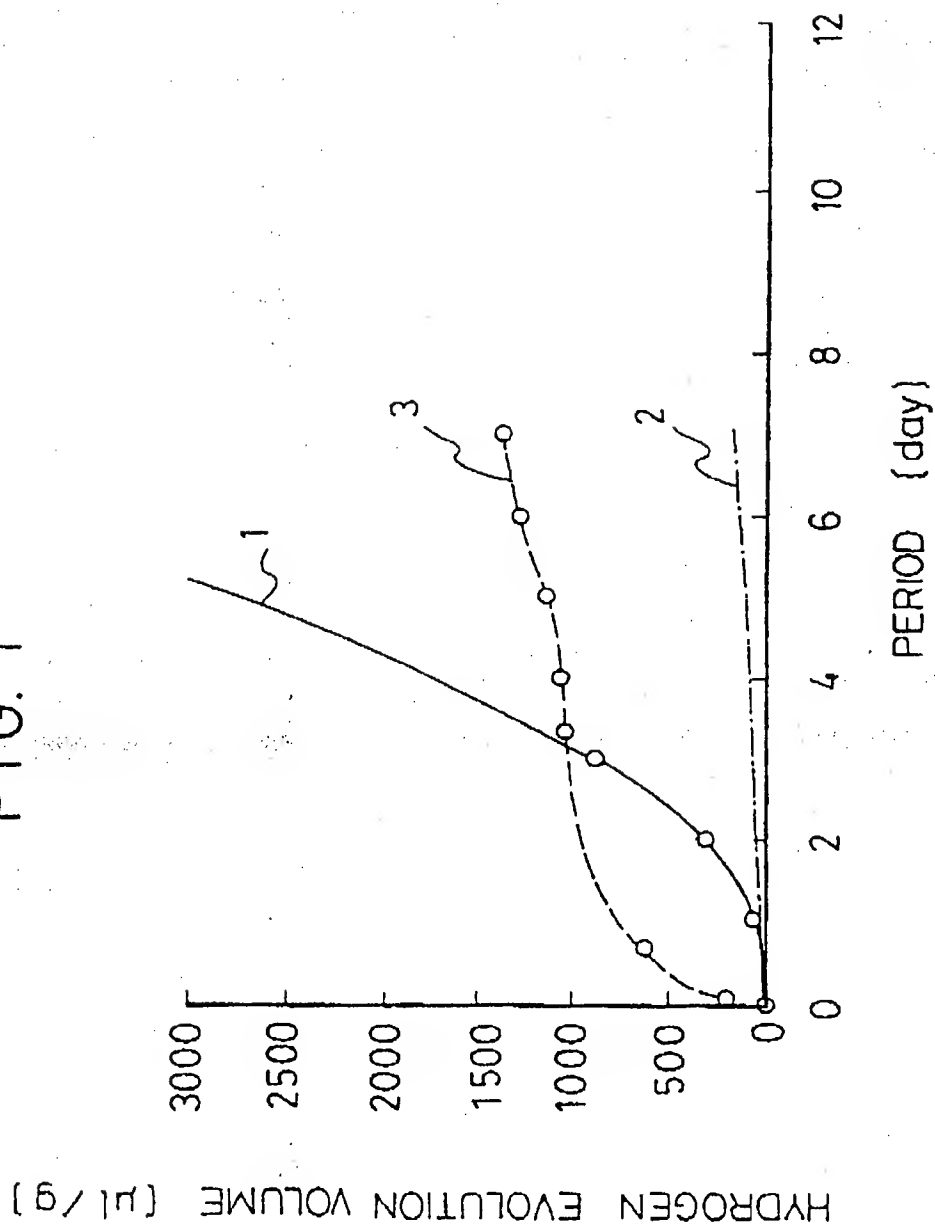
mercuryless zinc powder as a negative electrode active material, the negative electrode active material being added with two or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

44. An applied product using an alkaline battery comprising:

mercuryless zinc powder containing indium as a negative electrode active material; and
an electrolyte solution, the electrolyte solution being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.

45. An applied product using an alkaline battery comprising:
 mercuryless zinc powder containing indium as a negative electrode active material, the negative electrode active material being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.
46. An applied product using an alkaline battery comprising:
 mercuryless zinc powder as a negative electrode active material, a concentration of iron included in the mercuryless zinc powder being not more than 4 ppm with respect to zinc; and
 an electrolyte solution, the electrolyte solution being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.
47. An applied product using an alkaline battery comprising:
 mercuryless zinc powder as a negative electrode active material, a concentration of iron included in the mercuryless zinc powder being not more than 4 ppm with respect to zinc, the negative electrode active material being added with one or more species of compounds selected from an indium compound, a tin compound containing tetravalent tin, and lead oxide.
48. An applied product using a silver oxide battery comprising:
 mercuryless zinc as a negative electrode active material, the mercuryless zinc containing at least one species of metal selected from the group consisting of 0.001-0.5 % by weight of gallium, 0.005-0.5 % by weight of indium, 0.0001-0.5 % by weight of lead, 0.005-0.5 % by weight of bismuth, 0.001-0.5 % by weight of aluminum, and 0.001-0.5 % by weight of calcium, and the balance comprising zinc and inevitable impurities;
 an amount of water in the battery being 0.31-0.57 mg per a weight of 1 mg of the mercuryless zinc;
 and
 an electrolyte being potassium hydroxide.
49. An applied product using a silver oxide battery comprising:
 mercuryless zinc as a negative electrode active material, the mercuryless zinc containing at least one species of metal selected from the group consisting of 0.001-0.5 % by weight of gallium, 0.005-0.5 % by weight of indium, 0.0001-0.5 % by weight of lead, 0.005-0.5 % by weight of bismuth, 0.001-0.5 % by weight of aluminum, and 0.001-0.5 % by weight of calcium, and the balance comprising zinc and inevitable impurities;
 an amount of water in the battery being 0.32-0.59 mg per a weight of 1 mg of the mercuryless zinc;
 and
 an electrolyte being sodium hydroxide.
50. An applied product using an alkaline battery comprising:
 mercuryless zinc powder containing a gelling agent as a negative electrode active material;
 an outermost layer of a collector comprising a layer of zinc or metal having a hydrogen overvoltage higher than that of zinc; and
 an electrolyte solution, the electrolyte solution being added with one or more species of materials selected from an indium compound, lead oxide, hydroxide of alkaline earth metal, a surfactant having polyethylene oxide.
51. An applied product using an alkaline battery comprising:
 mercuryless zinc powder containing a gelling agent as a negative electrode active material, the negative electrode active material being added with one or more species of materials selected from an indium compound, lead oxide, hydroxide of alkaline earth metal, a surfactant having polyethylene oxide;
 and
 an outermost layer of a collector comprising a layer of zinc or metal having a hydrogen overvoltage higher than that of zinc.
52. An applied product using an alkaline battery comprising:
 mercuryless zinc powder as a negative electrode active material; and
 an outermost layer of a collector comprising an alloy layer containing zinc as an essential element, and containing one or more species selected from indium, lead and tin as a selective element.

FIG. 1



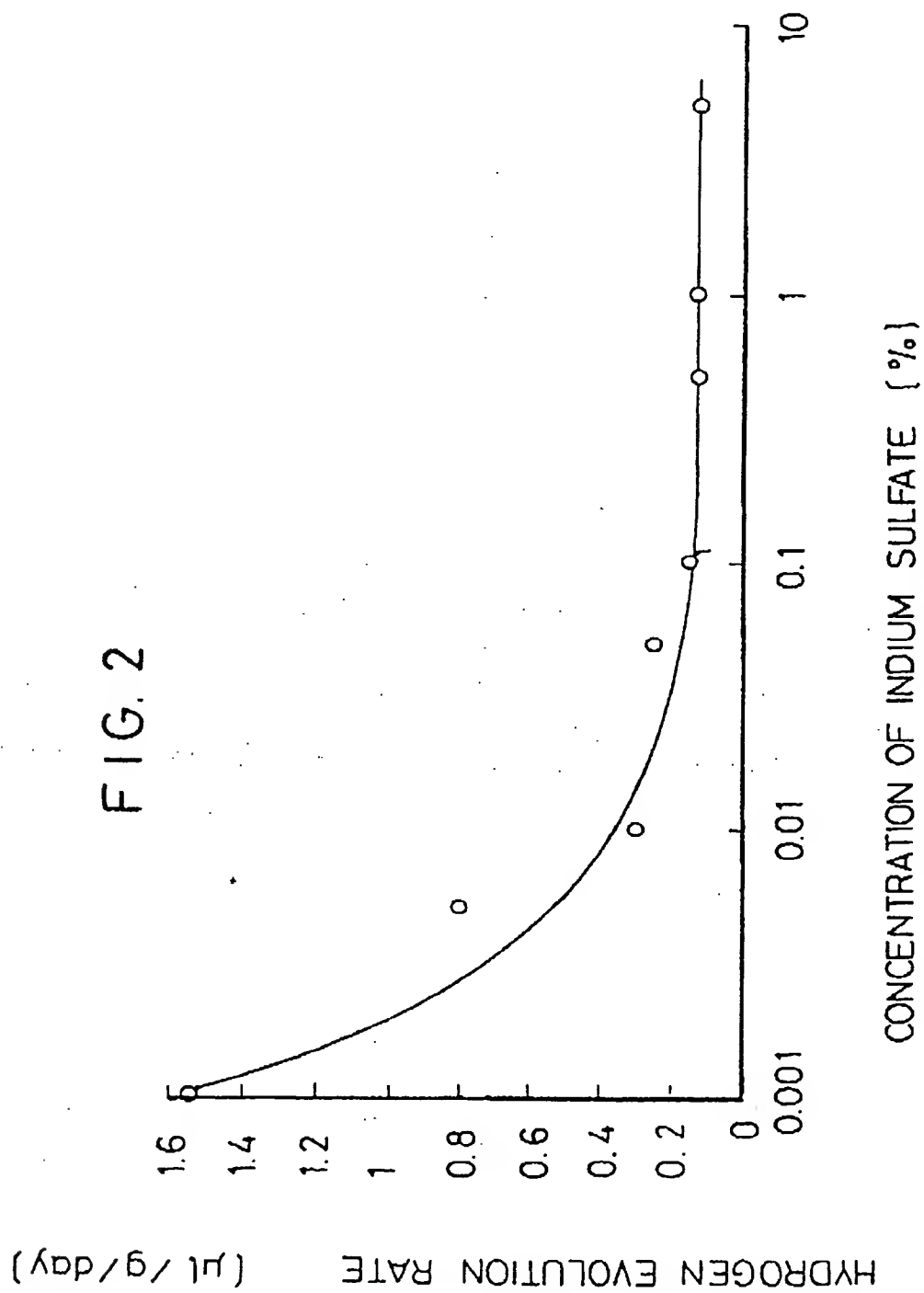


FIG. 3
ELECTRIC POTENTIAL VS. REFERENCE ELECTRODE (V)

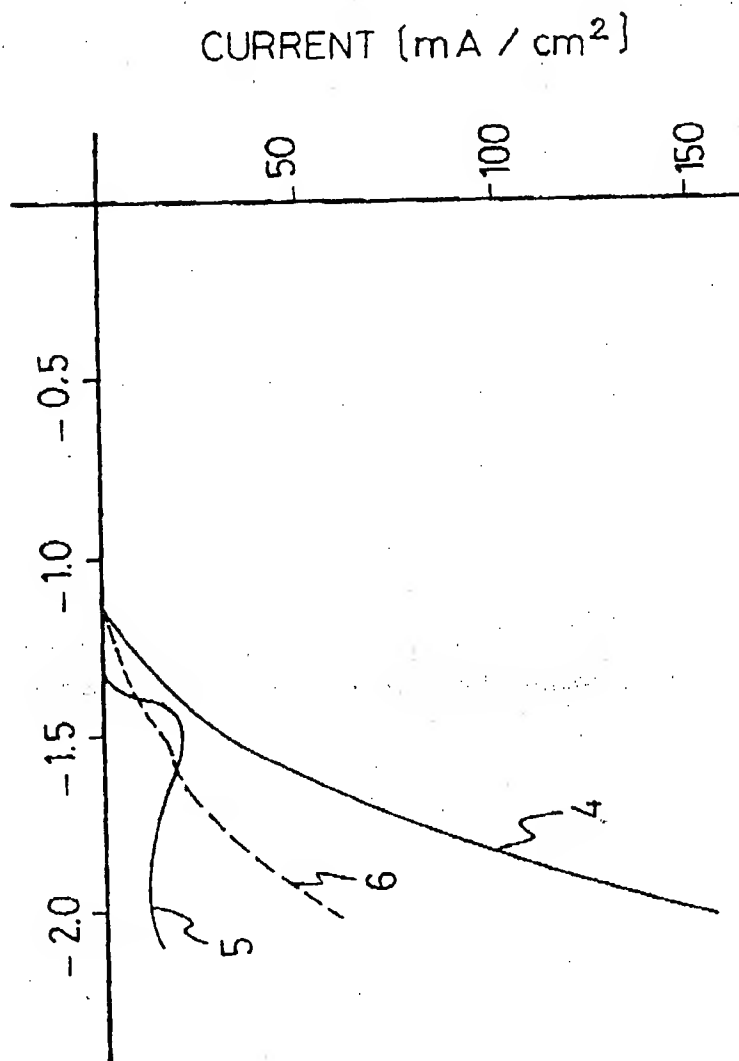


FIG. 4

ELECTRIC POTENTIAL VS REFERENCE ELECTRODE (V)

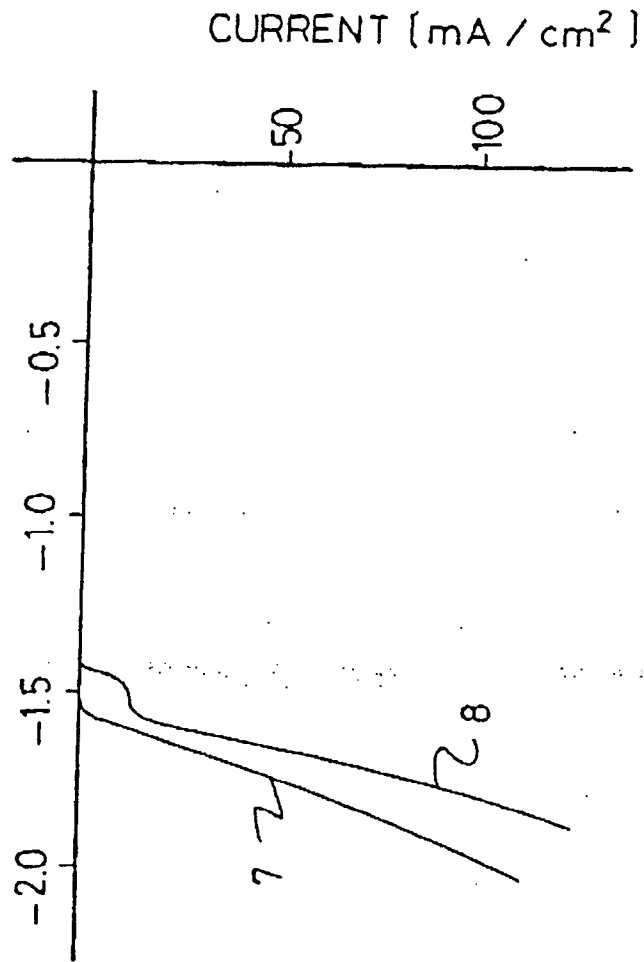
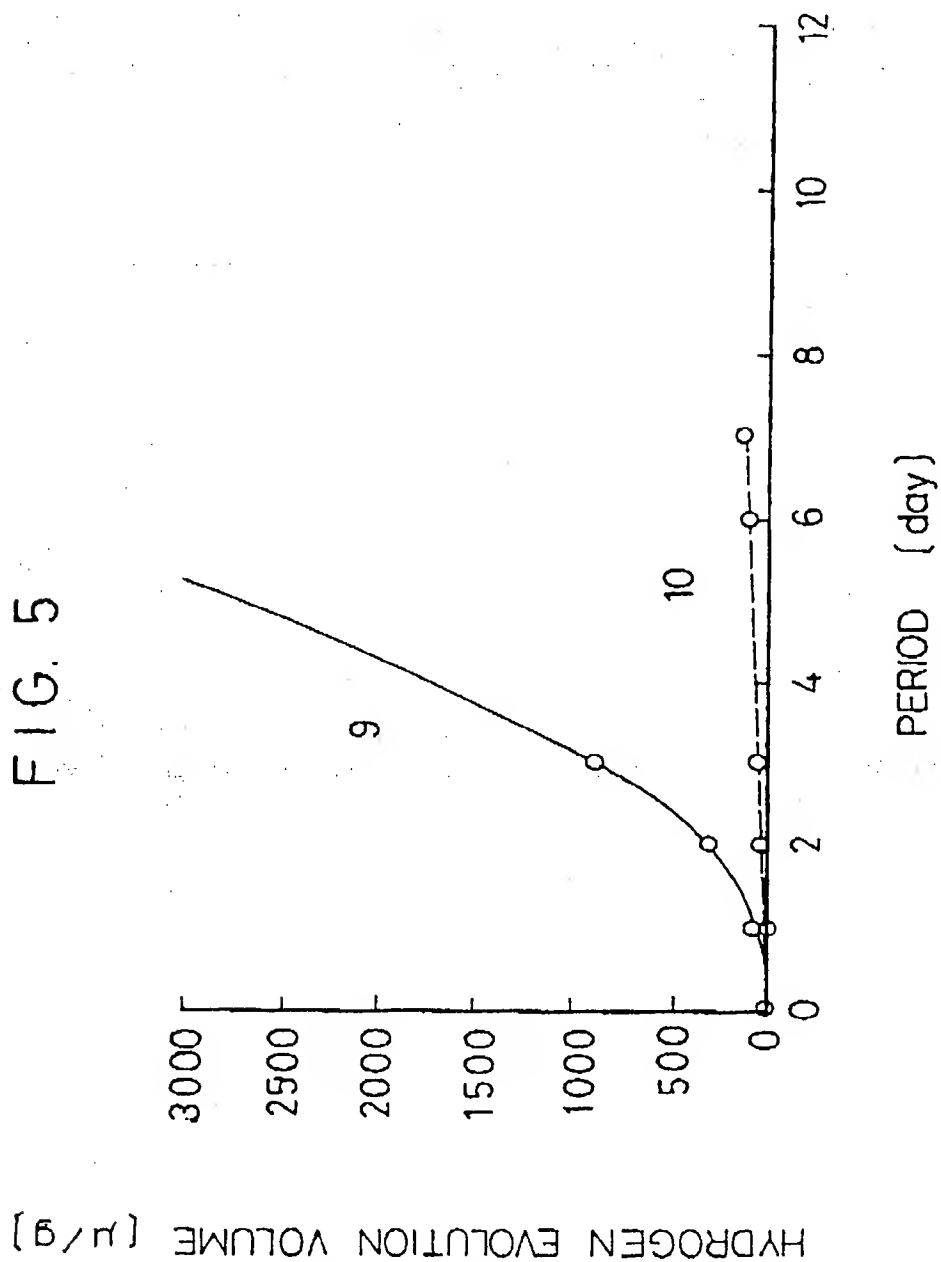


FIG. 5



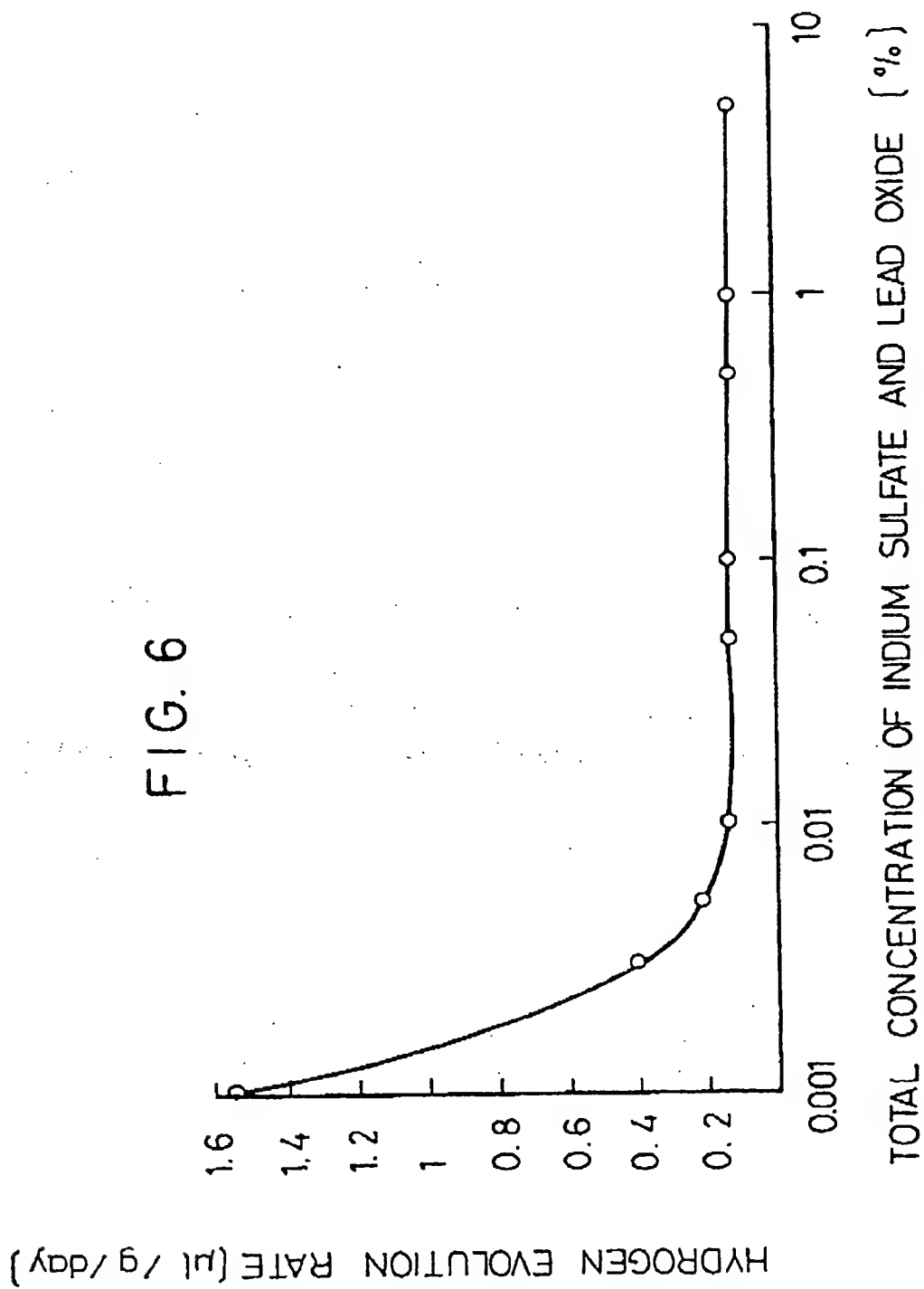
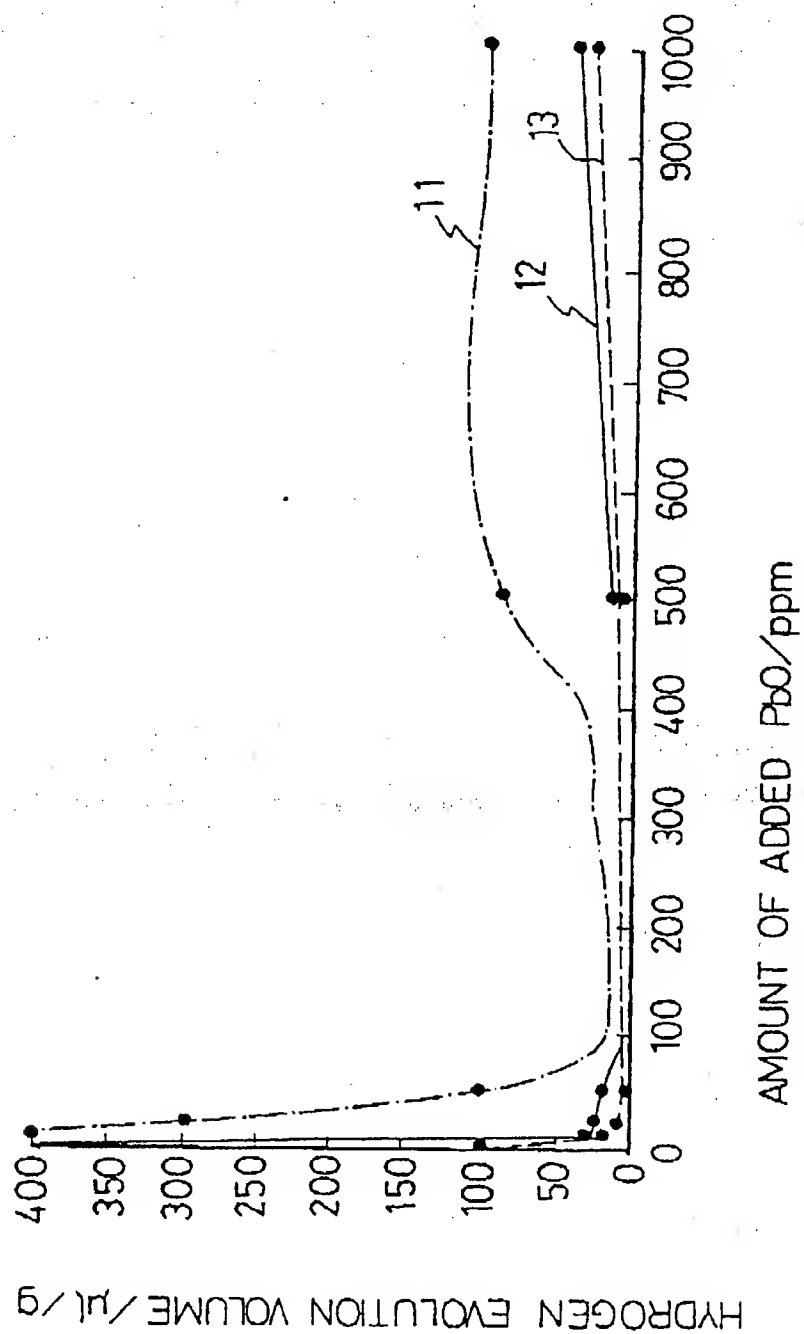
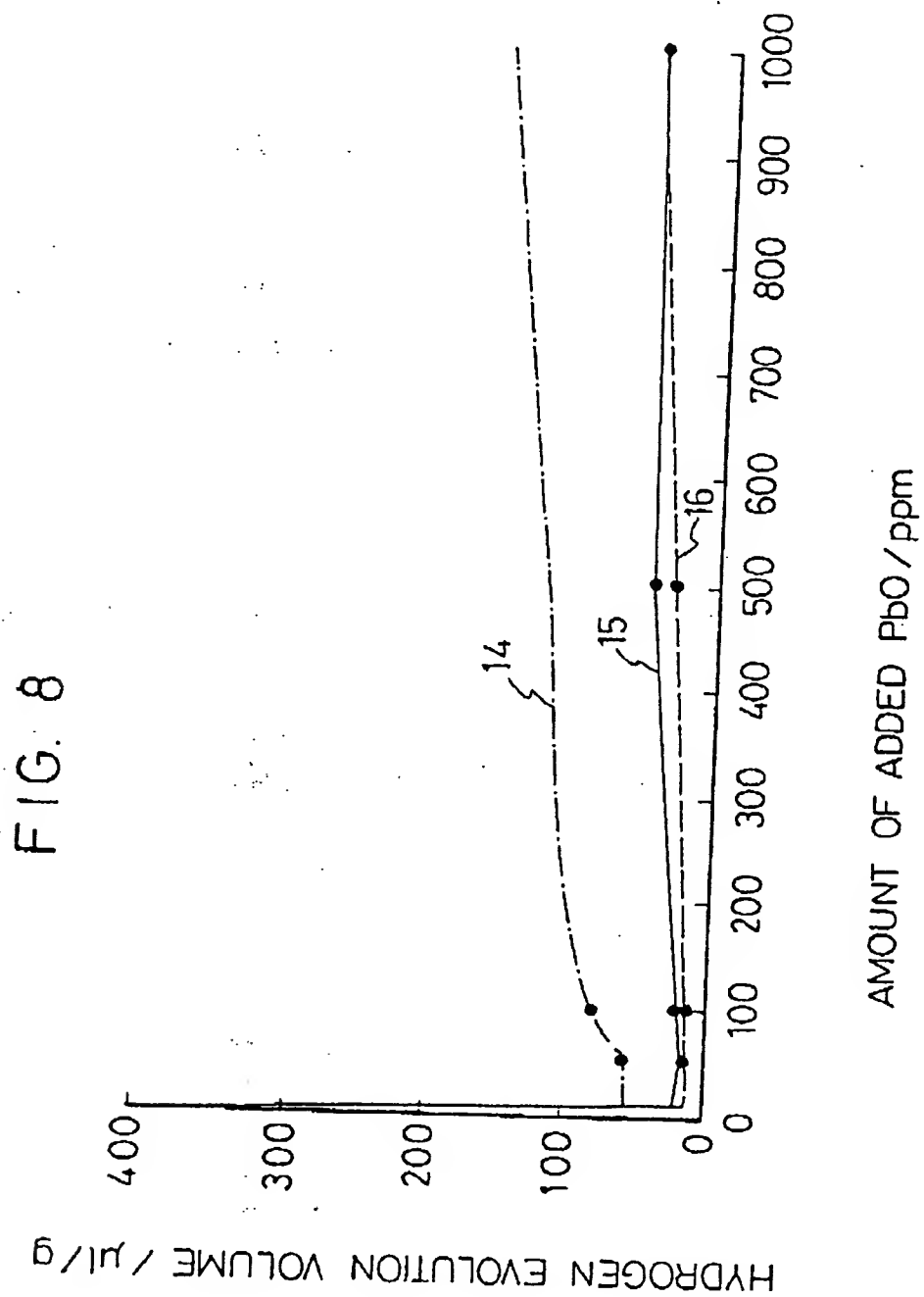


FIG. 7





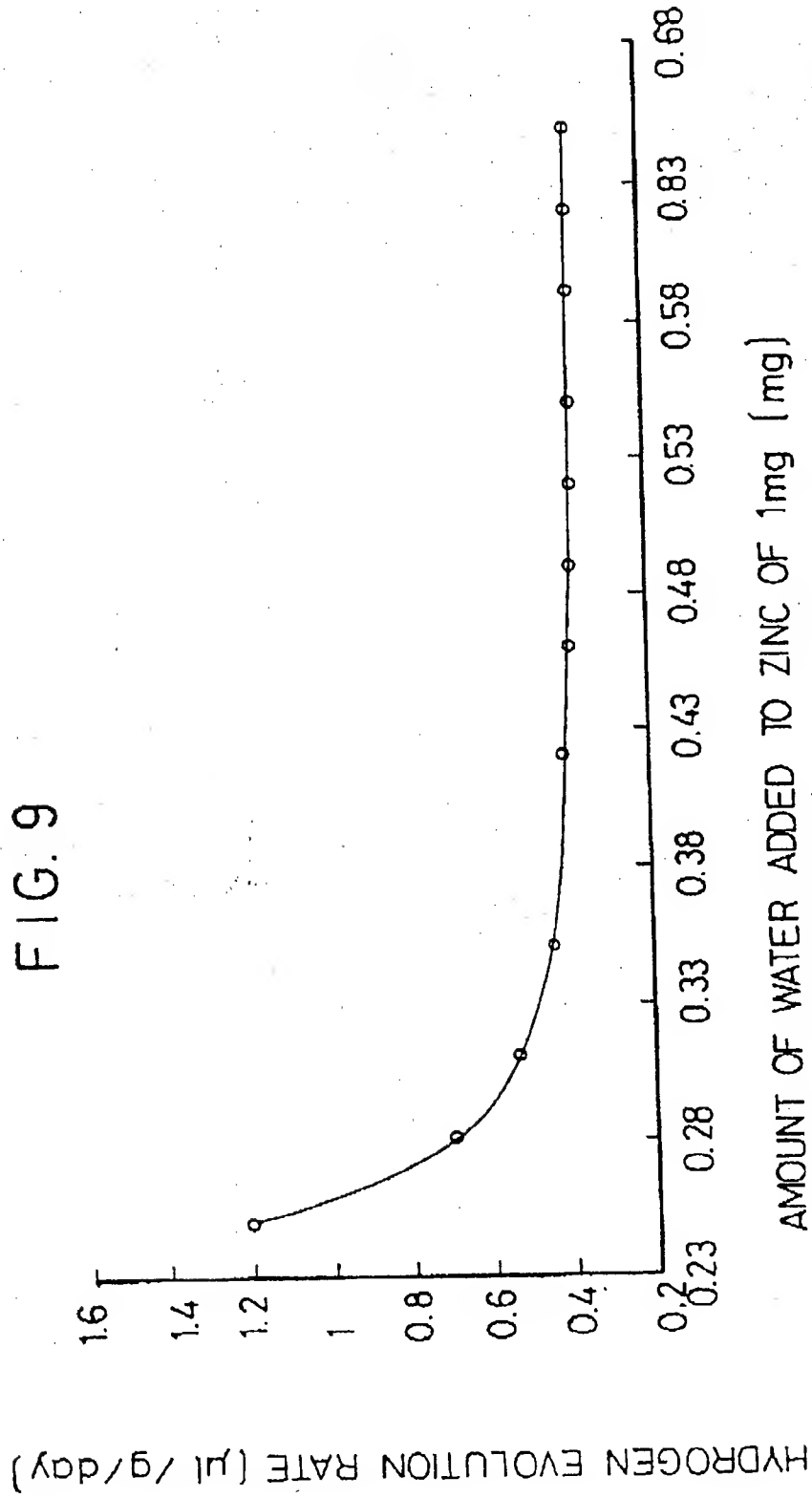
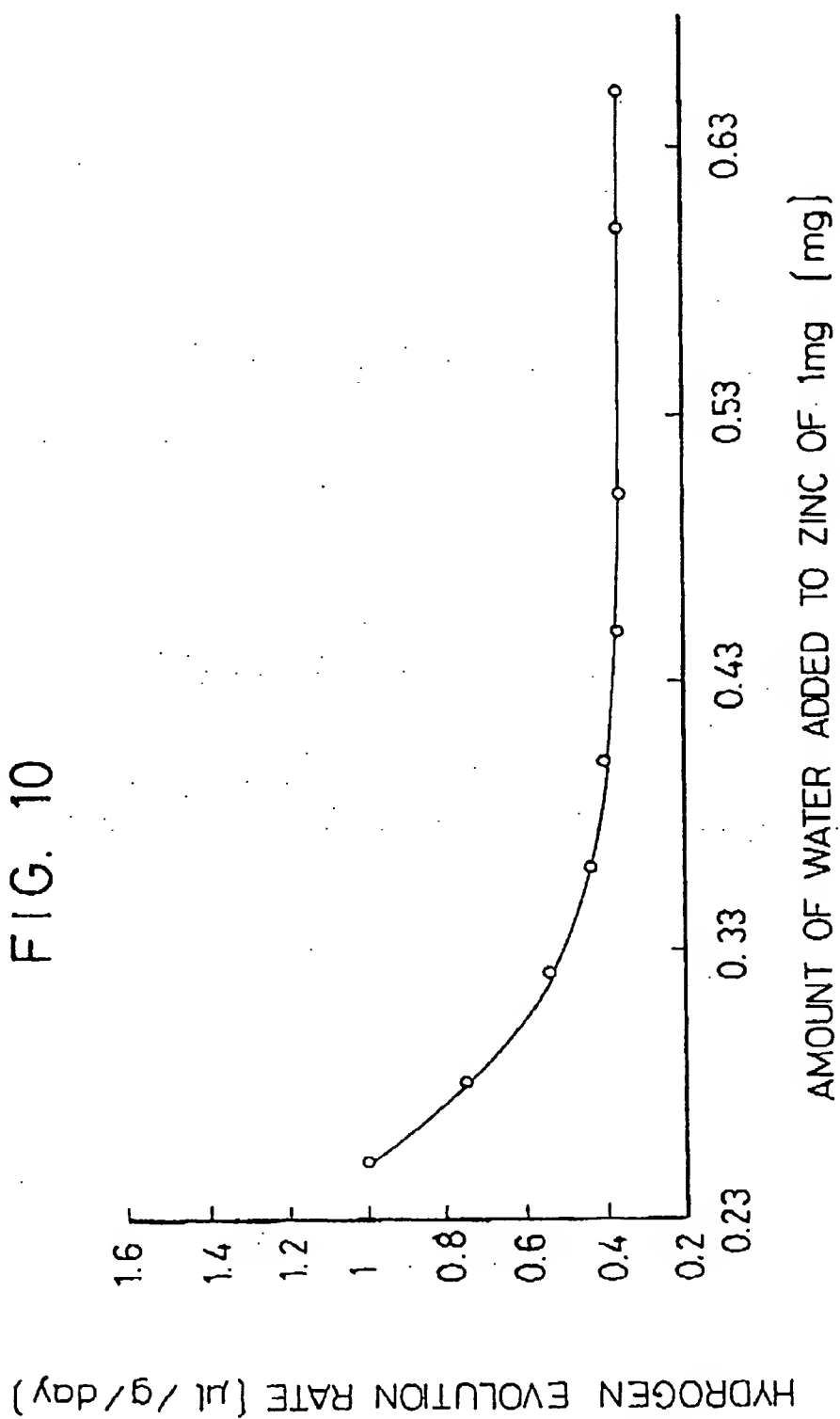


FIG. 10



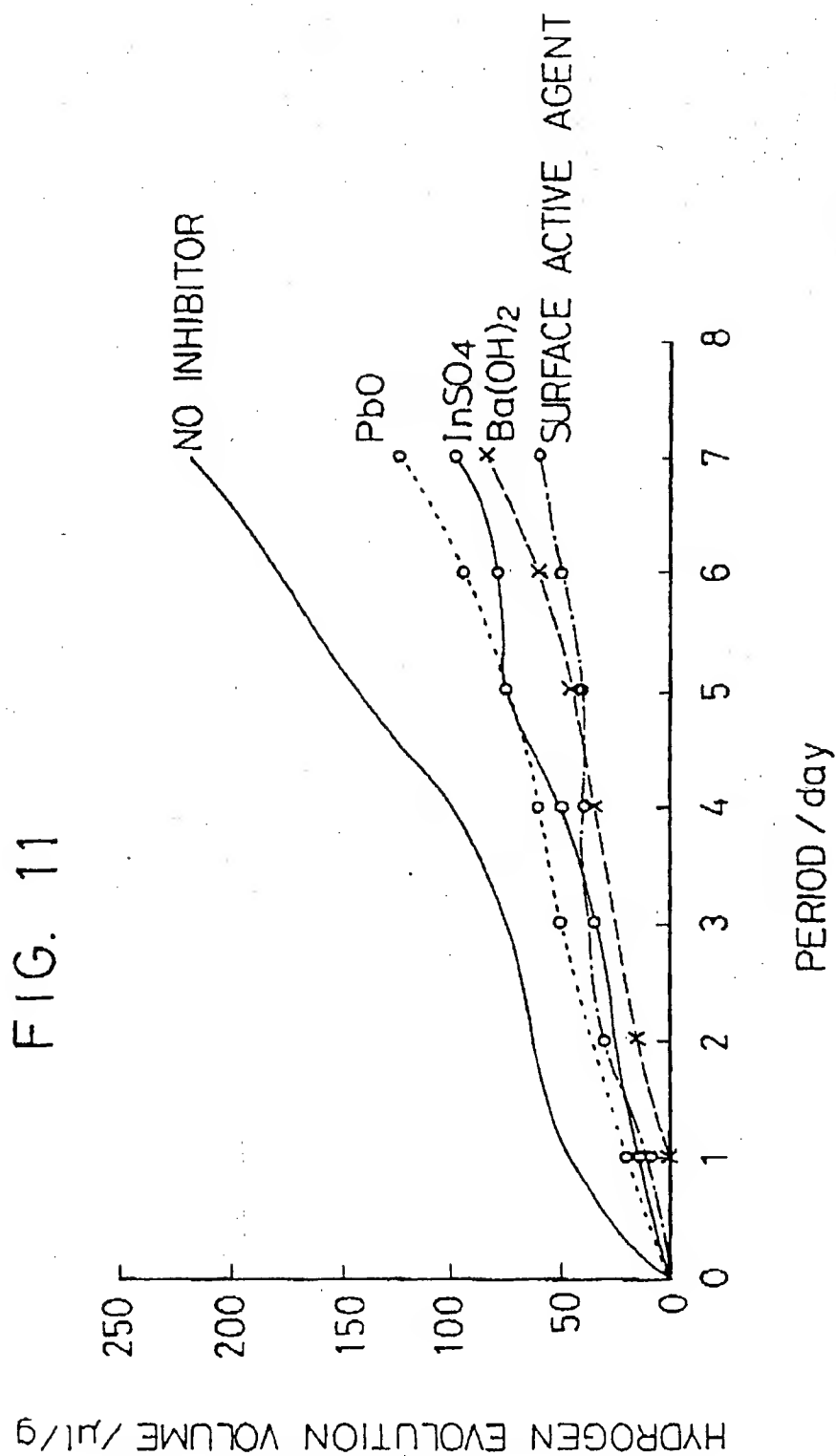


FIG. 12

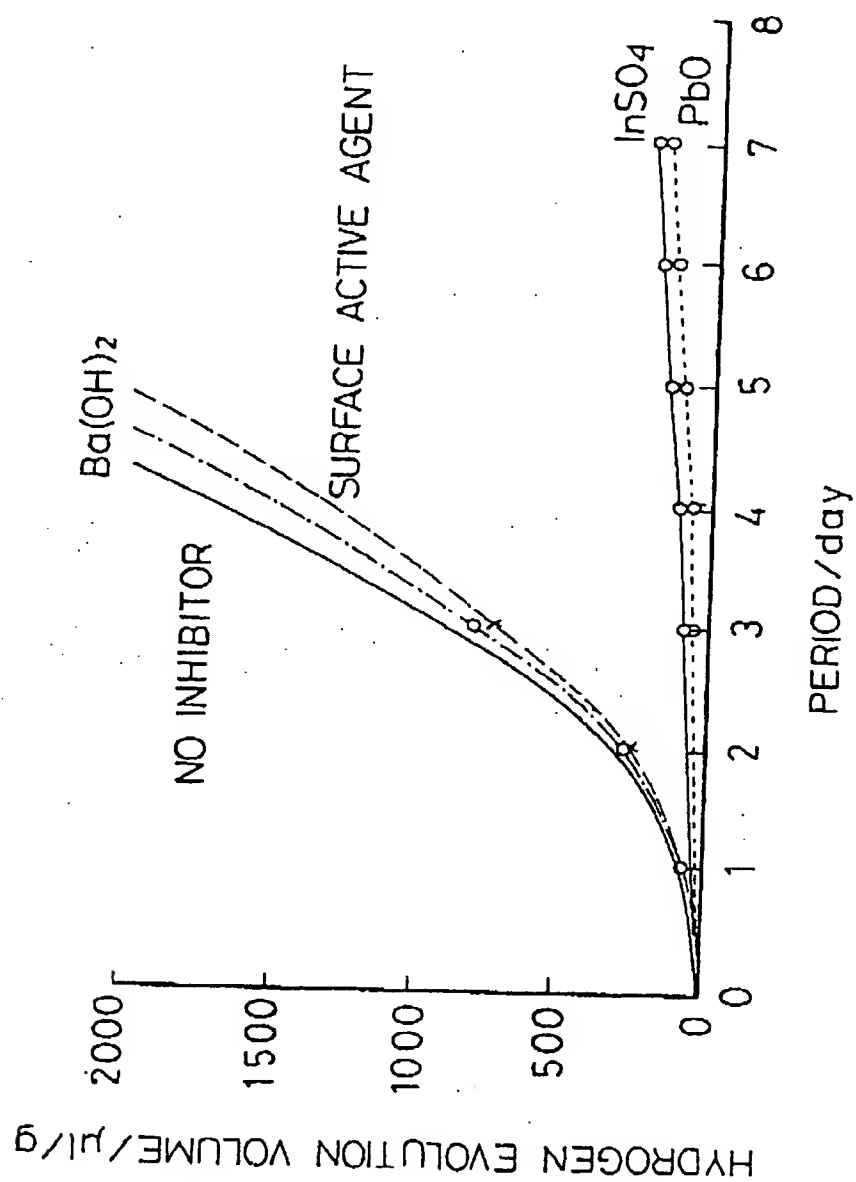
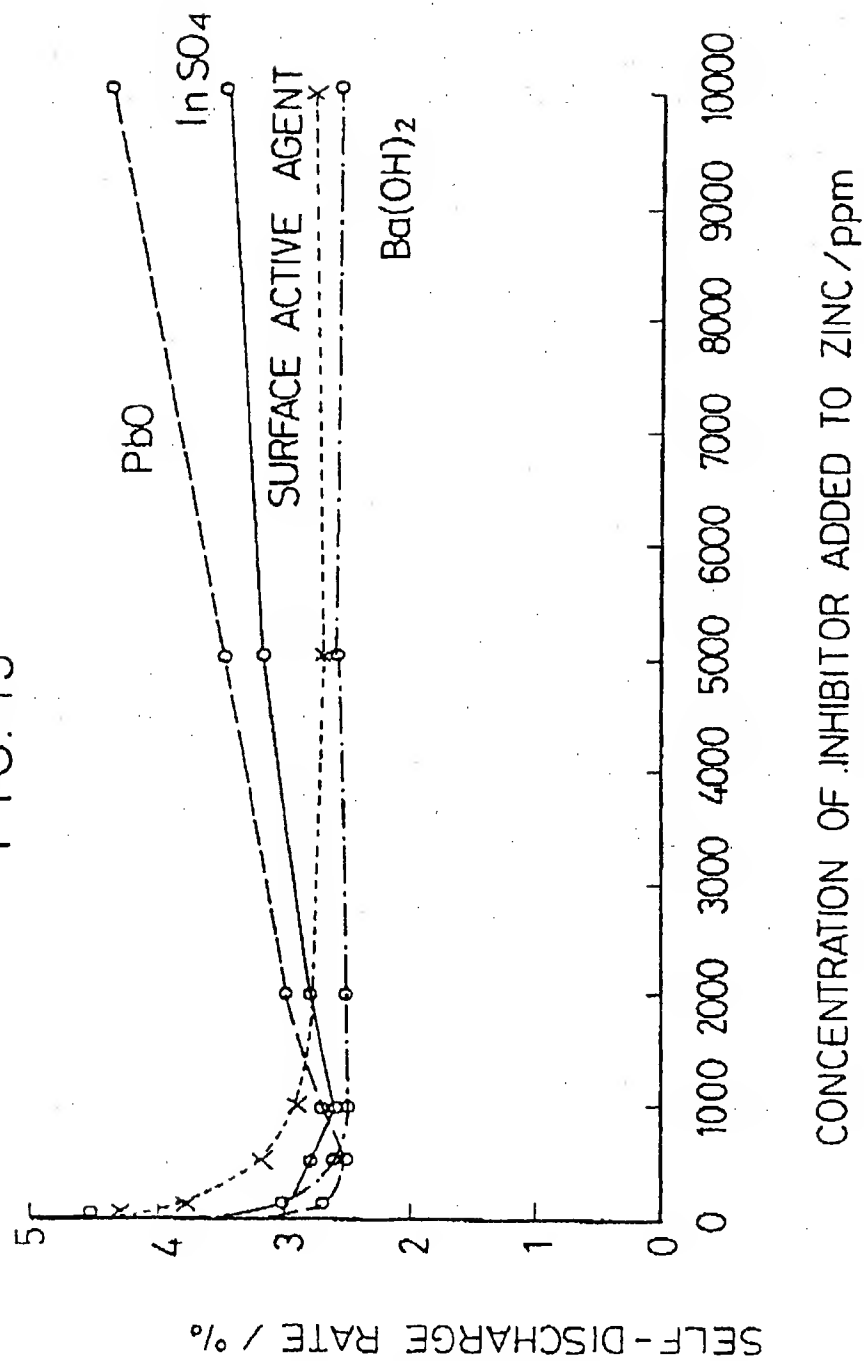
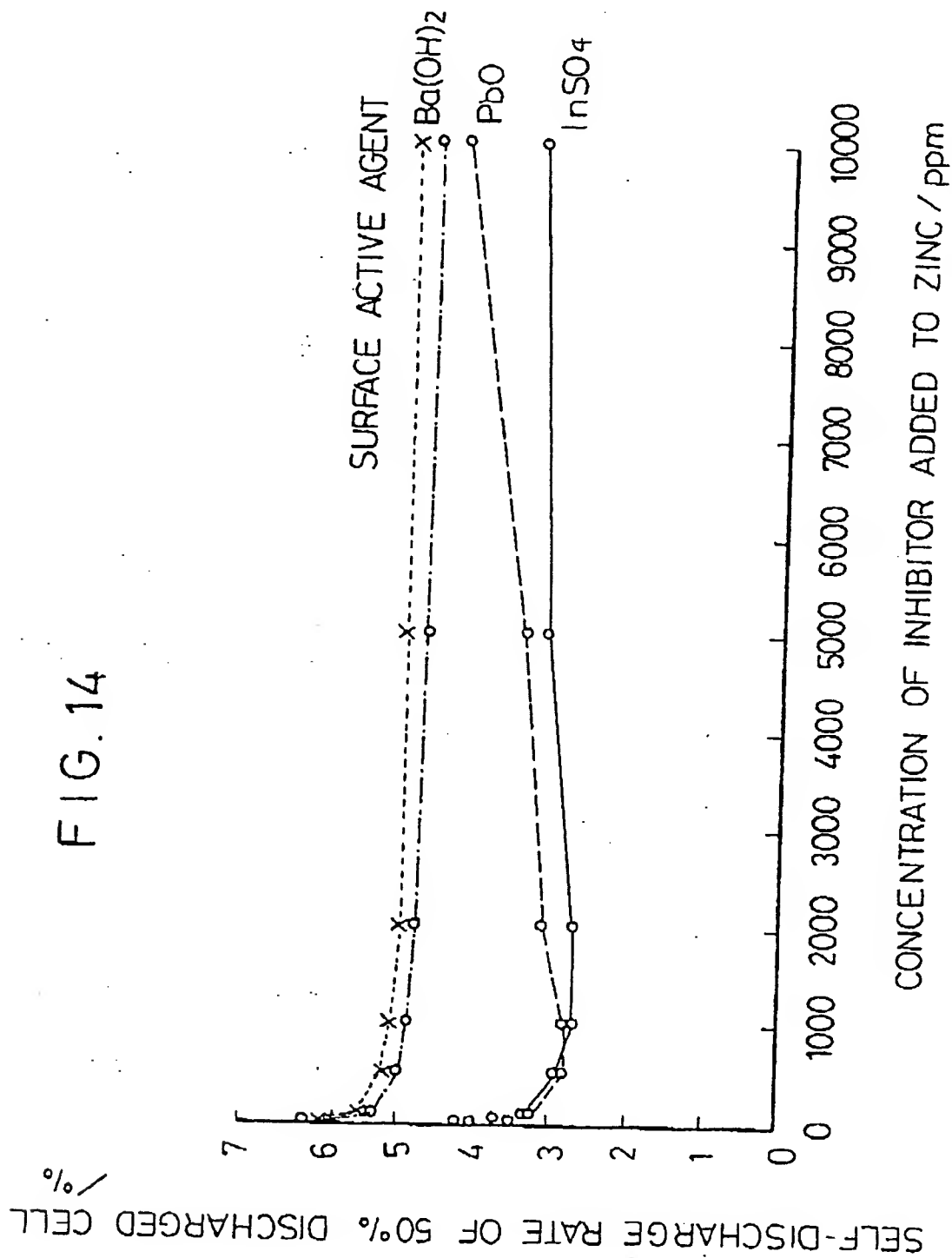


FIG. 13





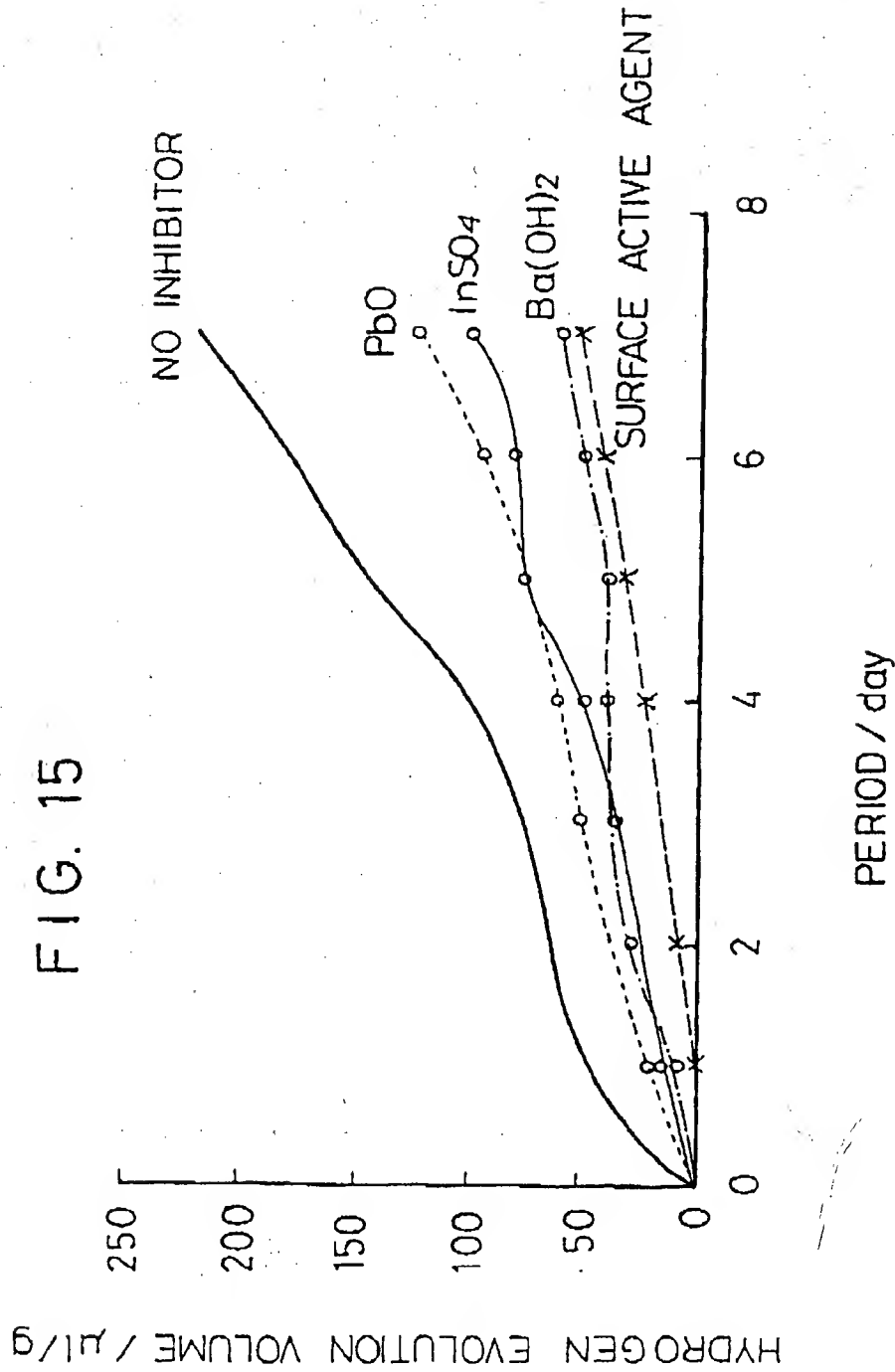
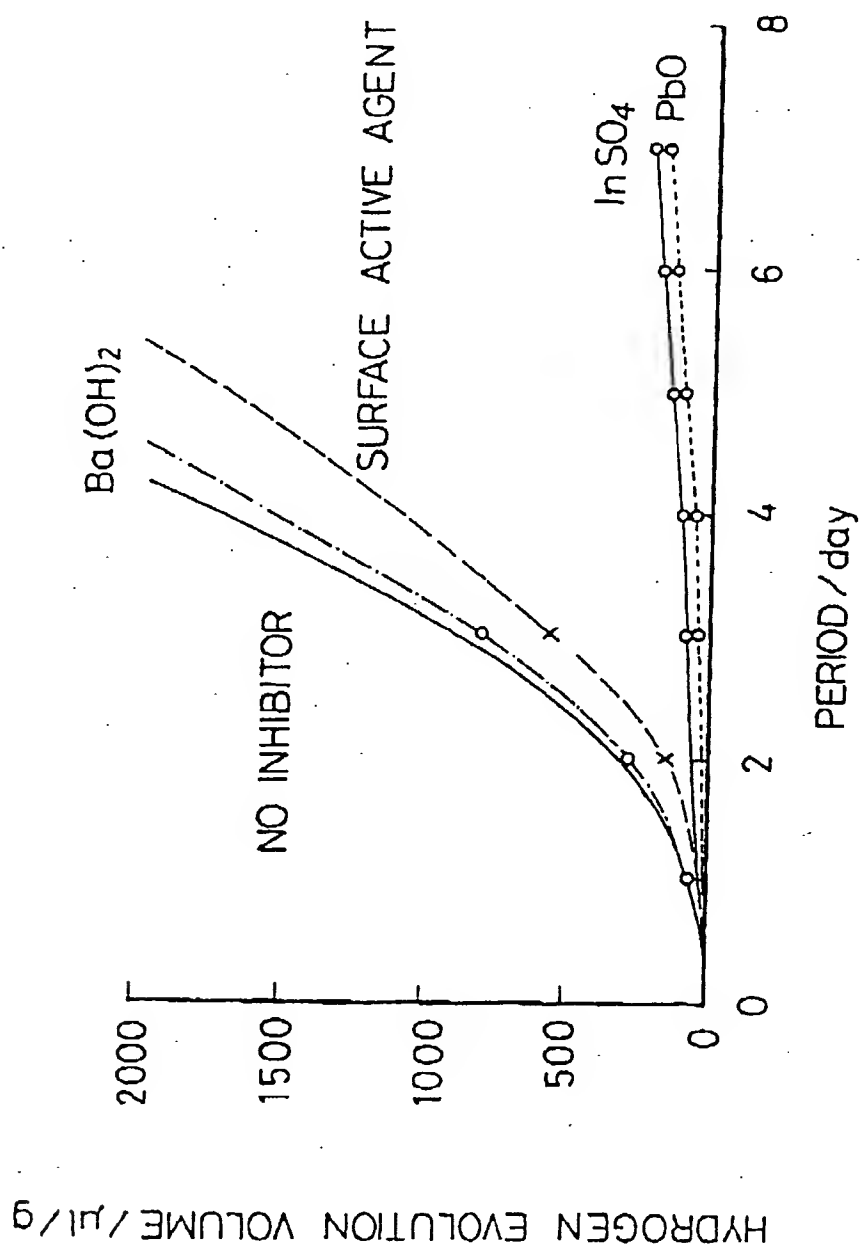
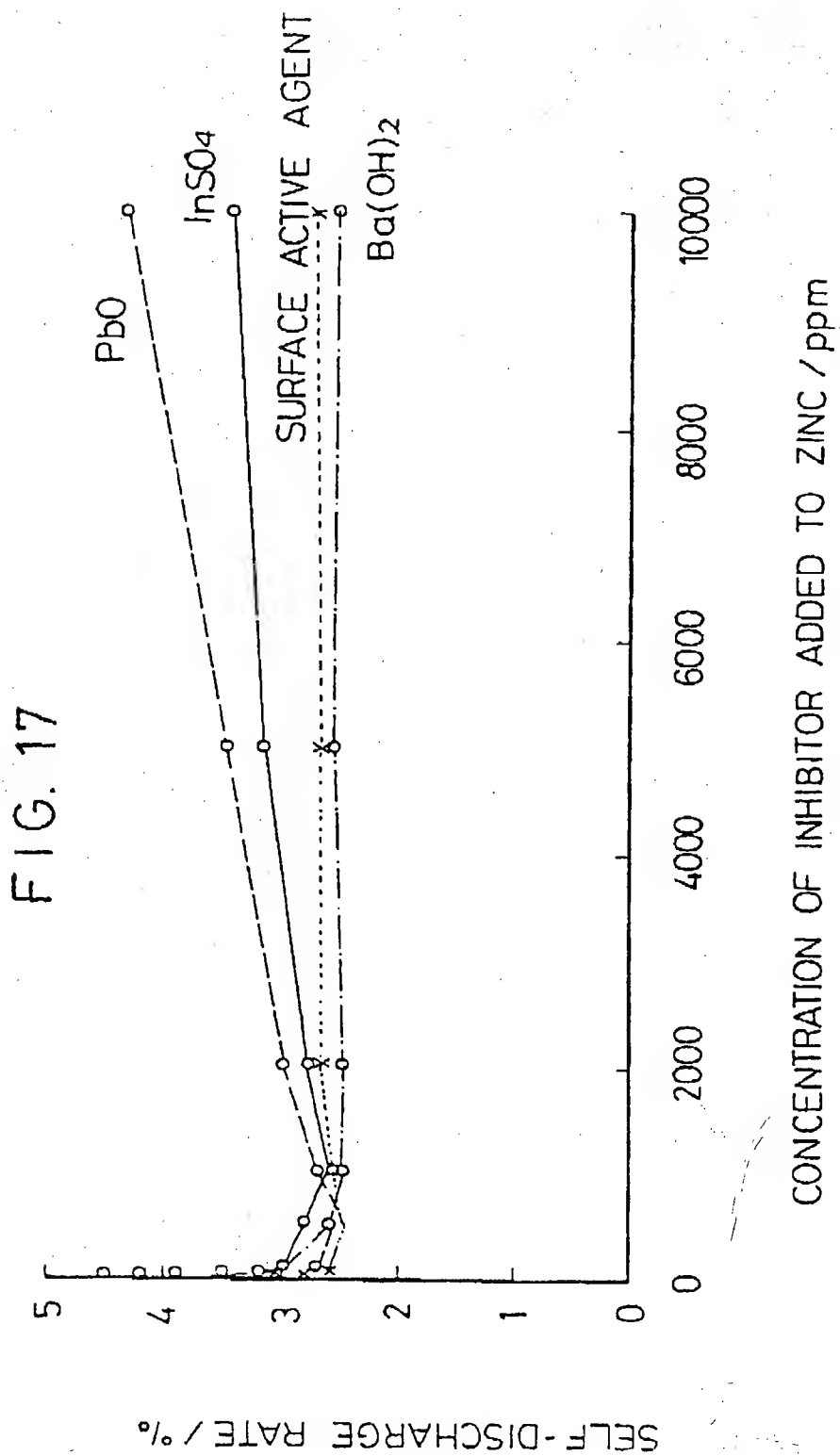


FIG. 16





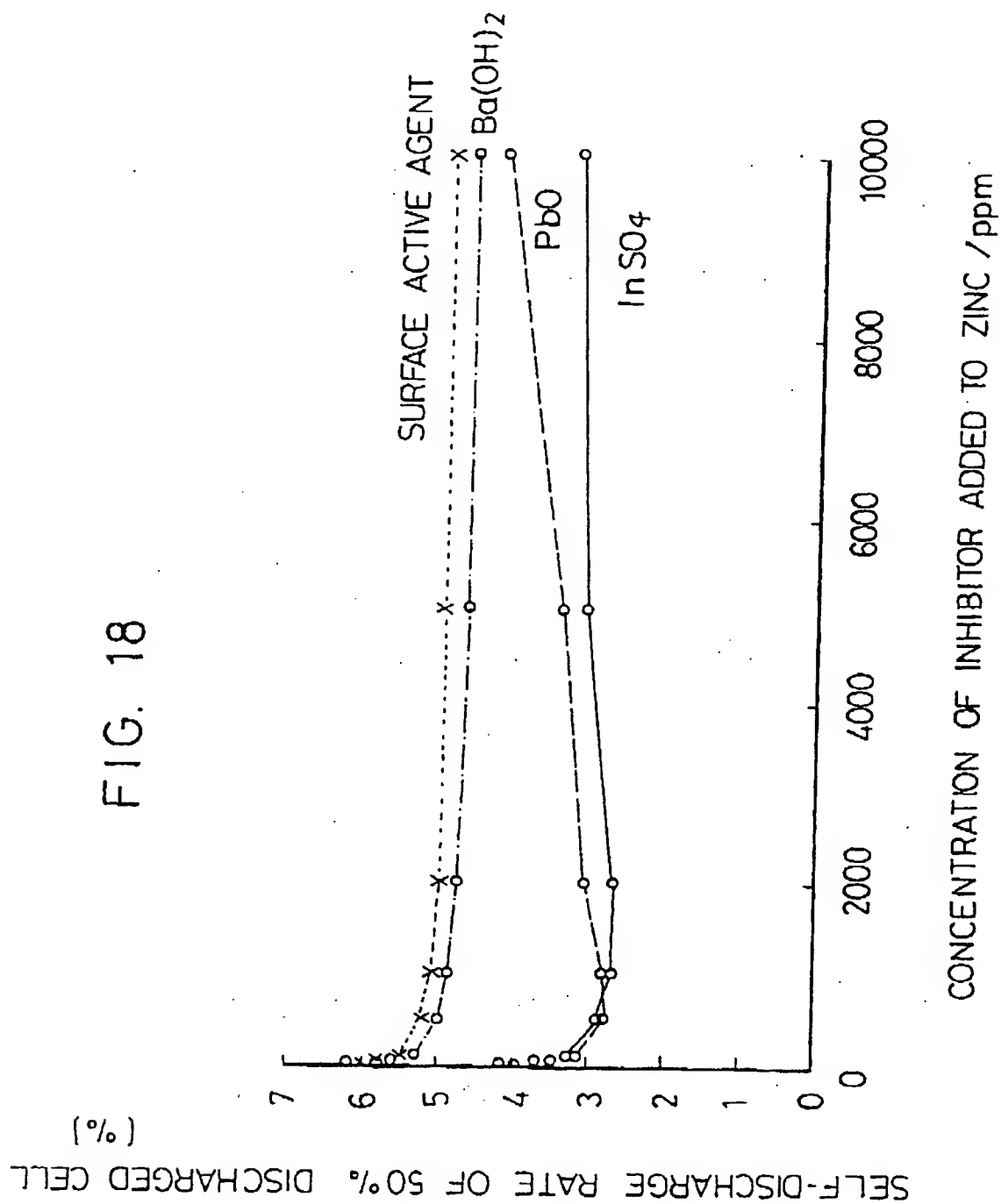
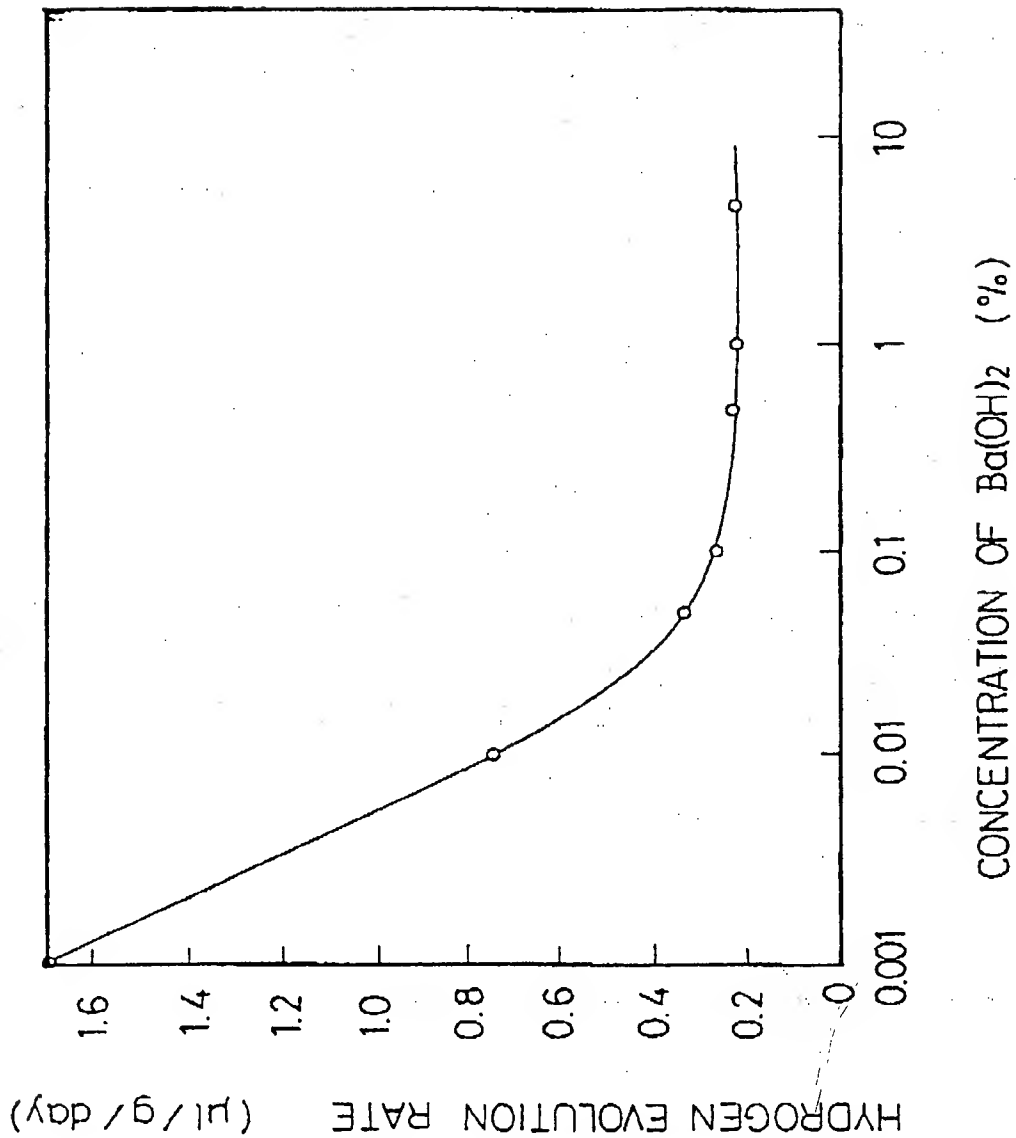


FIG. 19





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(71) Applicants:
• SEIKO INSTRUMENTS INC.
Tokyo 136 (JP)

• SEIKO ELECTRONIC COMPONENTS LTD.
Sendai-shi Miyagi (JP)

(72) Inventors:
• Watanabe, Shunji,
c/o Seiko Instruments Inc.
Tokyo (JP)
• Sakai, Tsuguo,
c/o Seiko Electronic Components Ltd
Sendai-shi, Miyagi (JP)
• Sakamoto Hideo
c/o Seiko Electronic Components Ltd
Sendai-shi, Miyagi (JP)

(74) Representative: Fleuchaus, Leo, Dipl.-Ing. et al
Melchiorstrasse 42
D-81479 München (DE)

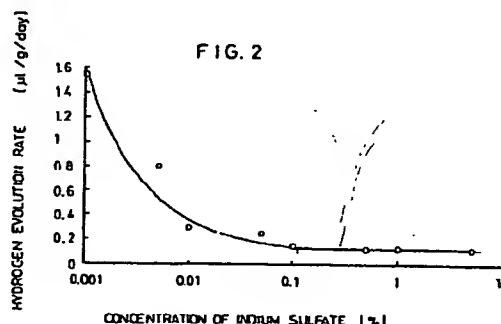
(54) Alkaline battery, production method of alkaline battery, and applied product using alkaline battery

(57) This invention relates to an alkaline battery in which hydrogen gas evolution and deterioration of battery performance can be prevented by using a collector having an outermost layer coated with zinc or metal having a higher hydrogen overvoltage than that of zinc and adding an inhibitor selected from an indium compound, lead oxide, a hydroxide of alkaline earth metal and polyoxyethylen alkylamide to an electrolyte solution or a negative electrode active material.

According to the present invention, it is possible to manufacture an alkaline battery in which evolution of hydrogen can be restrained without involving deterioration of battery performances. Especially, the present invention is effective in a coin type and a button type battery which is sensitive to the evolution of gas and is required to have high performance.

Further, the inventive collector which is plated with zinc alloy more than 3 μ m in thickness and used for low drain such as watches can attain a capacity equivalent to or more than that of a battery using amalgamated zinc. Further, watches loaded with the inventive alkaline batteries which are discarded do not pollute the environ-

ment, as they do not include mercury to be a pollution material.



EP 0 582 293 A3



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EUROPEAN SEARCH REPORT

Application Number
EP 93 11 2516

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.5)
X	J.ELECTROANAL. CHEM., vol. 180, no. 1-2, LAUSANNE, pages 495-510, XP000563463 L.BINDER: "CORROSION OF ZINC ELECTRODE MIXTURES IN ALKALINE MEDIA"	1,20,23, 24,27, 30,50,51	H01M6/06 H01M4/62 H01M4/66 H01M4/42 H01M4/12
Y	* page 501 - page 508, paragraph 4; table 1 * * page 495, paragraph 1 - page 496, paragraph 5 *	21,26, 28,33	
D,X	EP-A-0 457 354 (MATSUSHITA ELECTRIC INDUSTRIAL CO LTD)	18,19, 48,49	
Y	* page 3, line 24 - page 4, line 42; claims 1-4 * * page 7 - page 9; table 3 *	21,26, 28,33	
X	FR-A-2 634 594 (CIPEL) * page 5, line 28 - page 6, line 1; claims 1-3 *	2	
Y	EP-A-0 474 382 (EVEREADY BATTERY COMPANY) * claims 1-15 *	1	TECHNICAL FIELDS SEARCHED (Int.Cl.5)
X	* page 4, line 11 - page 6, line 6 *	16,17, 46,47	H01M
Y	GB-A-2 020 478 (ELECTROCHEM INC) * page 2, line 7-21 *	1	
X	DE-A-10 86 309 (DR. HANS MARTIN) * column 2, line 36 - column 3, line 22 * * column 6, line 4-63 *	1	
X	PATENT ABSTRACTS OF JAPAN vol. 5, no. 192 (E-085), 8 December 1981 & JP-A-56 116270 (SANYO ELECTRIC CO LTD) * abstract *	2	
The present search report has been drawn up for all claims			
Place of search		Date of completion of the search	Examiner
THE HAGUE		19 February 1996	DE VOS, A
CATEGORY OF CITED DOCUMENTS			
<p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 150 (03/82) (PM/C01)



EUROPEAN SEARCH REPORT

Application Number
EP 93 11 2516

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int. CL.5)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
X	FR-A-2 567 327 (DURACELL INTERNATIONAL INC) * page 3, line 15-33 *	2	
E	PATENT ABSTRACTS OF JAPAN vol. 18, no. 221 (E-1540), 20 April 1994 & JP-A-06 020674 (FUJI ELECTROCHEM CO LTD) * abstract *	1	
A	US-A-3 642 539 (AKIRA KAWAKAMI) * column 1, line 49 - column 2, line 6 *	1	
			TECHNICAL FIELDS SEARCHED (Int. CL.5)
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	19 February 1996	DE VOS, A	
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			



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CLAIMS INCURRING FEES

The present European patent application comprised at the time of filing more than ten claims.

- ☐ All claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for all claims.
- ☐ Only part of the claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been paid, namely claims:
- ☐ No claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims.

LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirement of unity of invention and relates to several inventions or groups of inventions, namely:

see sheet -B-

- ☐ All further search fees have been paid within the fixed time limit. The present European search report has been drawn up for all claims.
- ☐ Only part of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the inventions in respects of which search fees have been paid, namely claims:
- ☒ None of the further search fees has been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the invention first mentioned in the claims.

namely claims 1, 2, 16-33, 38, 39, 46-51



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EP 93 11 2516 -B-

LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirement of unity of invention and relates to several inventions or groups of inventions, namely:

1. Claims 1,2,16-33,38,39,46-51:

Alkaline battery with mercuryless zinc powder containing in electrolyte or negative electrode indium sulfate-sulfamate or -chloride as inhibitors. These compounds are used for their high solubility in alkaline solution (see page 34)

2. Claims 3-7,40,41:

Alkaline battery with mercuryless zinc powder as negative electrode containing in said negative electrode or in its electrolyte any indium compound in combination with a complexing agent for the indium ion. As less soluble known compounds, as indium hydroxide, are also include in these claims, they constitute a second inventive concept different from the invention of selection of indium compounds as defined in claims 1,2

3. Claims 8-13,42,43:

Alkaline battery with mercuryless zinc powder containing in the zinc anode or in the electrolyte a combination of two or more species selected from any indium compounds, tetravalent tin compound and lead oxide

4. Claims 14,15,44,45:

Alkaline battery with mercuryless zinc powder containing indium as negative electrode active material added with one or more species of any indium compounds, tetravalent in compounds or lead oxide added to electrolyte or negative electrode

5. Claims 34-37,52:

Alkaline battery with mercuryless zinc powder as negative active material and characterised by the use of a specific zinc alloy layer as outermost layer of the negative collector

